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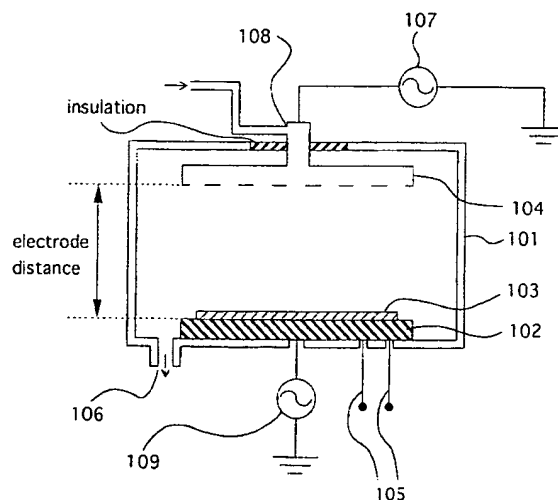
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(54) Film forming method and semiconductor device

(57) There is provided the film forming method of forming the insulating film 204 containing silicon on the

substrate 103 by plasmanizing the compound having the siloxane bonds and the oxidizing gas to react with each other.

FIG.1



Description

BACKGROUND OF THE INVENTION

1. Field of the Invention

[0001] The present invention relates to a method of forming an insulating film containing silicon, and to a semiconductor device and, more particularly to a technology valuable for forming the insulating film containing silicon having a low dielectric constant.

2. Description of the Prior Art

[0002] In recent years, miniaturization of the semiconductor device such as LSI, etc. progresses and thus wiring distances in the semiconductor device are reduced year by year. If the wiring distances are reduced in this manner, the parasitic capacitances of the wirings are increased and thus the operation speed of the semiconductor device is slowed down. Therefore, there is the possibility that the higher speed of the semiconductor device is prevented.

[0003] As one of the solutions to prevent such increase in the parasitic capacitances of the wirings, the insulating film having the low dielectric constant can be employed as the interlayer insulating film. The SiO_2 film is widely employed as the interlayer insulating film in the prior art. However, in order to reduce the wiring capacitances, the insulating film having the lower dielectric constant than this SiO_2 film must be employed. The relative dielectric constant of the SiO_2 film is 4.1, and up to the present there are following films that are known as the insulating film containing silicon having the lower dielectric constant than that of the SiO_2 film.

(1) SOG (Spin On Glass) film

(i) HSQ (Hydrogen Silsesquioxane) film Dielectric constant 3.1 to 3.5

(ii) MSQ (Methyl Silsesquioxane) film Dielectric constant 2.8 to 2.9

(2) FSG (Fluorinated Silicon Oxide) film Dielectric constant 3.5

[0004] With regard to the dielectric constant of above films, we referred to Table 1 set forth in "Monthly Semiconductor World", page 52, October, 1999.

SUMMARY OF THE INVENTION

[0005] It is an object of the present invention to provide a method of forming a noble insulating film containing silicon that is different from these insulating films in the prior art, and a semiconductor device using such insulating film.

[0006] According to the film forming method of the present invention, the insulating film containing silicon (hereinafter simply refer to as insulating film) is formed on the substrate by plasmanizing the compound having the siloxane bonds and the oxidizing gas to react with each other. Here the Si (silicon) atom in the "compound having the siloxane bonds" have already been bonded with O (oxygen) in the form of the siloxane bond (Si-O-Si). Therefore, in forming the film, it is difficult for C (carbon) to newly bond with the Si atom in the insulating film formed by using this compound. As a result, it is hard to form many Si-C bonds in the insulating film, and thus the number of Si-C bonds in the film is reduced.

[0007] It is generally known that the increase in the number of Si-C bonds in the film causes the increase in leakage current of the film. As described above, since the number of Si-C bonds in the film can be reduced in the insulating film formed by the film forming method of the present invention, the increase in the leakage current of the film can be suppressed.

[0008] Particularly, when H_2O is used as the oxidizing gas, an amount of NH_3 in the insulating film can be reduced. Therefore, when the chemical amplification resist coated on the insulating film is patterned, the resist is not crosslinked by NH_3 and thus the resist can be patterned finely. As a result, if the chemical amplification resist is used as an etching mask, it is possible to perform desirably the fine patterning of the insulating film underlying the resist.

[0009] Also, according to another film forming method of the present invention, the insulating film is formed on the substrate in the atmosphere in which the high-frequency power is applied to the upper and lower electrodes of the parallel plate type plasma chemical vapor deposition equipment, and the reaction gas containing $(\text{Si}(\text{CH}_3)_3)_2\text{O}$ and N_2O is introduced into the chamber. Here, $(\text{Si}(\text{CH}_3)_3)_2\text{O}$ is an example of the aforementioned "compound having the siloxane bond", and N_2O is an example of the aforementioned oxidizing gas. Therefore, as described above, the increase in the leakage current can be reduced in the insulating film formed by this film forming method.

[0010] Also, it becomes apparent that if the high-frequency power is applied to both the upper and lower electrodes

like this film forming method, the dynamic hardness of the insulating film can be increased compared to the case where the high-frequency power is applied only to the upper electrode.

[0011] In addition, according to the results of the examination made by the inventors of this application, it became apparent that if the pressure of the atmosphere in this case is set to more than 0.5 Torr, the dielectric constant of the insulating film can be reduced smaller than that of the conventional SiO_2 film.

[0012] Further, according to another film forming method of the present invention, the insulating film is formed on the substrate in the atmosphere in which the distance between the upper and lower electrodes of the parallel plate type plasma chemical vapor deposition equipment is set to more than 25 mm. In this case, the high-frequency power is applied only to the upper electrode and not to the lower electrode, and the reaction gas containing $(\text{Si}(\text{CH}_3)_3)_2\text{O}$ and N_2O is introduced into the chamber. As described above, the increase in the leakage current of the insulating film can be reduced. Besides, according to the results of the examination made by the inventors of this application, it became apparent that if the distance between the upper electrode and the lower electrode is set to more than 25 mm in this manner, the dielectric constant of the insulating film could be lowered.

[0013] Also, according to still another film forming method of the present invention, the $\text{Si}(\text{OR})_n\text{H}_m$ compound may be used in place of the compound having the siloxane bond. Here R denotes an alkyl group, $n+m=4$, and $m \geq 0$.

[0014] In this $\text{Si}(\text{OR})_n\text{H}_m$ compound, the Si atom is not directly bonded with R but bonded with R via O (oxygen). Hence, if the $\text{Si}(\text{OR})_n\text{H}_m$ compound is used, Si-C bonds are difficult to be formed in the insulating film. Therefore, like the compound having the siloxane bonds, it is difficult to form a large number of Si-C bonds in the insulating film by using the $\text{Si}(\text{OR})_n\text{H}_m$ compound, and thus the number of Si-C bonds in the film can be reduced. As a result, the increase in the leakage current due to many Si-C bonds can be reduced in this insulating film.

[0015] It should be noted that a plasma process may be employed after the formation of the insulating film, which is formed using the compound having the siloxane bonds or the $\text{Si}(\text{OR})_n\text{H}_m$ compound, in order to improve the hygroscopicity resistance of the film. In this plasma process, the atmosphere containing at least one of O_2 , N_2O , and NH_3 is plasmanized and then the surface of the insulating film is exposed to the plasmanized atmosphere.

[0016] According to this, H_2O contained in the film and CO_2 formed in this plasma process by oxidizing C in the film are discharged to the outside of the film, and also unbonded bonds of Si in the film are terminated by O, N, H, etc. Therefore, unbonded bonds of Si in the insulating film can be prevented from being bonded by OH group and the like, and thus the hygroscopicity resistance of the insulating film can be improved.

BRIEF DESCRIPTION OF THE DRAWINGS

[0017]

FIG. 1 is a sectional view showing a plasma CVD (Chemical Vapor Deposition) equipment used to form an insulating film containing silicon according to the present invention;

FIGS. 2A and 2B are sectional views showing a method of forming the insulating film containing silicon according to an embodiment of the present invention;

FIG. 3 is a graph showing change of high-frequency powers applied in forming the insulating film containing silicon according to the embodiment of the present invention with respect to a time;

FIG. 4 is a sectional view showing a method of forming the insulating film containing silicon according to an embodiment of the present invention when an underlying insulating film and a cover insulating film are formed;

FIG. 5 is a graph showing a relationship between a relative dielectric constant of an insulating film 204 containing silicon and a pressure and a relationship between a refractive index and the pressure when a pressure of the atmosphere is changed while applying a high-frequency power only to an upper electrode 104 by setting an electrode distance to 25 mm;

FIG. 6 is a graph showing a relationship between the relative dielectric constant of the insulating film 204 containing silicon and the pressure and a relationship between the refractive index and the pressure when the pressure of the atmosphere is changed while applying the high-frequency power to both the upper electrode 104 and a lower electrode 102 by setting the electrode distance to 25 mm;

FIG. 7 is a graph showing a relationship between a power of a second high-frequency power supply 109 and a dynamic hardness of the insulating film 204 containing silicon and a relationship between the power and a Young's modulus of the insulating film 204 containing silicon when a power of the second high-frequency power supply 109 is changed by setting the electrode distance to 25 mm;

FIG. 8 is a graph showing a relationship between a power of a first high-frequency power supply 107 and the relative dielectric constant of the insulating film 204 containing silicon and a relationship between the power and a refractive index of the insulating film 204 containing silicon when a power of the first high-frequency power supply 107 is changed by setting the electrode distance to 50 mm;

FIG. 9 is a graph showing a relationship between the power of the second high-frequency power supply 109 and

the relative dielectric constant of the insulating film 204 containing silicon and a relationship between the power and the refractive index of the insulating film 204 containing silicon when the power of the second high-frequency power supply 109 is changed by setting the electrode distance to 50 mm;

FIG. 10 is a graph showing a relationship between the relative dielectric constant of the insulating film 204 containing silicon and the pressure and a relationship between the refractive index and the pressure when the pressure of the atmosphere is changed while using H_2O as an oxidizing gas by setting the electrode distance to 50 mm;

FIG. 11 is a graph showing results of the insulating film 204 containing silicon measured by the Infrared Absorption Spectroscopy when the pressure of the atmosphere is changed while using H_2O as the oxidizing gas by setting the electrode distance to 25 mm;

FIG. 12 is a graph showing a relationship between the power of the second high-frequency power supply 109 and the relative dielectric constant of the insulating film 204 containing silicon and a relationship between the power and the refractive index of the insulating film 204 containing silicon when the power of the second high-frequency power supply 109 is changed while using H_2O as the oxidizing gas by setting the electrode distance to 25 mm;

FIG. 13 is a sectional view used to explain a method of measuring a leakage current in the insulating film 204 containing silicon;

FIG. 14 is a graph showing characteristics of the leakage current in the insulating film 204 containing silicon when the power of the second high-frequency power supply 109 is changed while using H_2O as the oxidizing gas by setting the electrode distance to 25 mm;

FIG. 15 is a graph showing measured results of an amount of NH_3 , that is contained in an insulating film formed by using $Si(CH_3)_4$ and N_2O , measured by the TDS method;

FIG. 16 is a graph showing measured results of an amount of NH_3 , that is contained in the insulating film 204 containing silicon formed by using HMDS and H_2O , measured by the TDS method;

FIG. 17 is a graph showing a relationship between the pressure of the atmosphere and the relative dielectric constant of the insulating film 204 containing silicon and a relationship between the pressure and the refractive index of the insulating film 204 containing silicon when the pressure of the atmosphere is changed while using HMDS and H_2O ;

FIG. 18 is a graph showing characteristics of the leakage current in the insulating film 204 containing silicon when the pressure of the atmosphere is changed while using HMDS and H_2O ;

FIGS. 19A to 19F are sectional views showing a sectional shape respectively when the damascene process is applied to the insulating film 204 containing silicon according to the embodiment of the present invention; and

FIGS. 20A to 20G are sectional views showing a sectional shape respectively when the damascene process is applied to the insulating film 204 containing silicon formed by using H_2O as the oxidizing gas according to the embodiment of the present invention.

DESCRIPTION OF THE PREFERRED EMBODIMENT

[0018] An embodiment of the present invention will be explained in detail with reference to the accompanying drawings hereinafter.

(1) Explanation of the plasma CVD equipment used to form an insulating film in the present invention.

[0019] FIG. 1 is a sectional view showing the plasma CVD (Chemical Vapor Deposition) equipment used to form the insulating film containing silicon (hereinafter simply refer to as insulating film) in the present embodiment.

[0020] In FIG. 1, a reference 101 denotes a chamber used to form the film, and two opposing electrodes, i.e., a lower electrode 102 and an upper electrode 104 are provided in the chamber 101. Here, the lower electrode 102 serves also as a loading table on which a substrate 103 is loaded. A heater (not shown) for heating the substrate 103 up to the desired temperature is built in the lower electrode 102. A reference 105 denotes power supplying lines that supply a power to this heater.

[0021] Further, the upper electrode 104 serves also as a shower head that supplies a gas to an interior of the chamber 101.

[0022] A first high-frequency power supply 107 and a second high-frequency power supply 109 are connected to these two electrodes 104 and 102 respectively. The gas in the chamber 101 can be plasmanized by supplying a high-frequency power to the gas in the chamber 101 from one of these high-frequency power supplies 107, 109, or from both the power supplies.

[0023] In addition, a gas introducing port 108 is provided to the upper electrode 104, and the gas is introduced into the chamber 101 via the gas introducing port 108. An exhaust port 106 is provided to the chamber 101, and the gas introduced into the chamber 101 is exhausted via the exhaust port 106 to reduce a pressure in the chamber 101.

[0024] It should be noted that since the upper electrode 104 and the lower electrode 102 are provided in the chamber

101 to oppose to each other, this plasma CVD equipment is so-called parallel plate type plasma chemical vapor deposition equipment.

(2) Explanation of the method of forming the insulating film according to the present invention.

(i) Outline

[0025] The inventors of this application have made following consideration before making the present invention.

[0026] First, it is considered that in order to form the insulating film having the low dielectric constant, the SiO_2 film should be formed to contain groups containing C (carbon) and H (hydrogen) in the film so as to reduce a density of the SiO_2 film. It is expected that when the SiO_2 film is formed in this manner, Si-O bonds are disconnected at the portion where these groups exist and thus the dielectric constant of the portion is reduced, which in turn result in reducing the dielectric constant of the overall film.

[0027] Second, it is considered that when the SiO_2 film is formed as above, the above groups should be contained in the film not to form a large number of Si-C bonds. This is because, if a large number of Si-C bonds are formed in the film, portions in which a large number of Si-C bonds are continuously exist are formed in the film, and thus there is the possibility that the leakage currents are generated in these portions.

[0028] Third, it is considered that, in view of the first and second considerations, ① a reaction gas containing the compound having the siloxane bond (Si-O-Si) or ② a reaction gas containing the $\text{Si}(\text{OR})_n\text{H}_m$ compound should be employed as the reaction gas used to form the film. Since the Si-O bonds are formed originally in these compounds, there is the possibility that it can be made difficult to form a large number of Si-C bonds when forming the film.

[0029] Then, examples of these compounds will be listed hereunder specifically.

① Compound having the siloxane bond (Si-O-Si)

HMDS (hexamethyldisiloxane: $(\text{Si}(\text{CH}_3)_3)_2\text{O}$)

OMCTS (octomethylcyclotetrasiloxane: $(\text{Si}(\text{CH}_3)_2)_4\text{O}_4$)

BEDS (hexaethyldisiloxane: $(\text{Si}(\text{C}_2\text{H}_5)_3)_2\text{O}$)

TMDS (tetramethyldisiloxane: $(\text{SiH}(\text{CH}_3)_2)_2\text{O}$)

TEDS (tetraethyldisiloxane: $(\text{SiH}(\text{C}_2\text{H}_5)_2)_2\text{O}$)

TMCTS (tetramethylcyclotetrasiloxane: $(\text{SiH}(\text{CH}_3))_4\text{O}_4$)

TETCS (tetraethylcyclotetrasiloxane: $(\text{SiH}(\text{C}_2\text{H}_5))_4\text{O}_4$)

[0030] These compounds are liquid at the room temperature (20 °C).

② $\text{Si}(\text{OR})_n\text{H}_m$ compound

$\text{Si}(\text{OCH}_3)_3\text{H}$

$\text{Si}(\text{OC}_2\text{H}_5)_3\text{H}$

$\text{Si}(\text{OCH}_3)_4$

$\text{Si}(\text{OC}_2\text{H}_5)_4$

[0031] These compounds are a liquid at the room temperature (20 °C). It should be noted that the $\text{Si}(\text{OR})_n\text{H}_m$ compounds are not limited to these specific examples, but may use such a compounds satisfying $n+m=4$ and $m \geq 0$, where n and m denotes integer.

(ii) Explanation of film forming conditions

[0032] Next, film forming conditions will be explained with reference to FIGS.2A and 2B, when the above compounds are employed to form the insulating film containing silicon. FIGS.2A and 2B are sectional views showing a method of forming the insulating film according to an embodiment of the present invention.

[0033] At first, as shown in FIG.2A, a silicon substrate 201 on a surface of which a BPSG (borophosphosilicate glass) film 202 is formed is prepared. Then, an aluminum film is formed on the BPSG film 202 and then aluminum wirings 203 are formed by patterning the aluminum film. These silicon substrate 201, the BPSG film 202, and the aluminum wirings 203 constitute the substrate 103.

[0034] Then, as shown in FIG.2B, an insulating film 204 containing silicon (simply referred to as insulating film hereinafter) is formed on the substrate 103. This is performed by introducing the reaction gas via the gas introducing port 108 after the substrate 103 is loaded on the lower electrode 102 in the chamber 101 (see FIG.1), then heating the lower electrode 102 up to the predetermined temperature, and then applying the high-frequency power to the reaction gas from the first and second high-frequency power supplies 107, 109. At the same time, an inside of the chamber

101 is set to a predetermined pressure by exhausting the reaction gas via the exhaust port 106.

[0035] As described above, as compounds contained in the reaction gas, there are ①the compound having the siloxane bond (Si-O-Si) and ②the $\text{Si}(\text{OR})_n\text{H}_m$ compound. Two cases where these compounds are employed respectively will be explained in the following.

①the case where the compound having the siloxane bond (Si-O-Si) is used

[0036] The film forming conditions in this case will be given in Table 1.

Table 1

Flow rate of compound having siloxane bond (Si-O-Si)	50 sccm
Flow rate of an oxidizing gas	see Table 2
Flow rate of an inert gas (Ar or He)	200 sccm
Temperature of a substrate 103	350 °C
Pressure in a chamber 101	1.75 Torr
Frequency of the high frequency power applied by the first high frequency power supply 107	13.56 MHz
Power of the high frequency power applied by the first high frequency power supply 107	50 to 300 W
Frequency of the high frequency power applied by the second high frequency power supply 109	380 KHz
Power of the high frequency power applied by the second high frequency power supply 109	10 to 100 W

[0037] As the compound having the siloxane bond (Si-O-Si) set forth in the first line of Table 1, there are HMDS, OMCTS, HEDS, TMDS, TEDS, TMCTS, and TECTS, which are already mentioned above. Any one of these compounds may be contained in the reaction gas. These compounds, which are a liquid at the room temperature, are heated to be vaporized after the flow rates of them are controlled by the liquid mass flowmeter (not shown), and then introduced into the chamber 101. Alternatively, the flow rates of these compounds may be controlled by the high-temperature mass flowmeter (not shown) after these compounds are vaporized, and then these compounds may be supplied to the chamber 101. The flow rate of the compound having the siloxane bonds given in Table 1 is the flow rate of the compound vaporized in these manners.

[0038] Also, there are N_2O , O_2 , H_2O , CO_2 as the oxidizing gas set forth in the second line of Table 1, and at least one of them may be contained in the reaction gas. In case only one of these oxidizing gases is added into the reaction gas, i.e., without combination of other oxidizing gases, the flow rate of the oxidizing gas will be given in Table 2.

Table 2

Oxidizing gas	Flow rate
N_2O	50 to 200 sccm
O_2	10 to 50 sccm
H_2O	100 to 300 sccm
CO_2	100 to 300 sccm

[0039] The inert gas set forth in the third line of Table 1 is not essential. That is, even if such inert gas is not contained in the reaction gas, advantages similar to those described in the following can be achieved.

[0040] The dielectric constant of the insulating film 204 formed in accordance with these conditions was 2.7 at 1 MHz. This value is smaller than that of the conventional SiO_2 film (4.1).

[0041] The reason why the insulating film 204 having such a low dielectric constant can be formed is considered as follows. That is, as can be seen from their chemical formulas, the compounds having the siloxane bond (Si-O-Si) contain the alkyl groups such as CH_3 , C_2H_5 and the like. Therefore, if the insulating film 204 is formed using these compounds, the alkyl groups are contained in the film. In the insulating film 204, the Si-O bonds are disconnected at the portions where the alkyl groups are contained, which in turn creates pores at the portions and reduce the dielectric constant of that portions. As a result, the dielectric constant of the overall insulating film 204 is lowered.

[0042] In addition, Si and O have already been bonded in the compound having the siloxane bond (Si-O-Si). Therefore, since the bonding of Si and C is limited in the insulating film 204, the Si-C bonds that cause the increase in the leakage current in the film are hard to be formed. Accordingly, it is expected that the leakage current of the insulating

film 204 can be reduced compared with the case where the compound that does not have the siloxane bond (Si-O-Si) is used.

[0043] Though the high-frequency power applied by the first and second high-frequency power supply 107 and 109 are constant in time in the film forming conditions given in Table 1 and Table 2, high-frequency power whose power varies in time as shown in FIG.3 may also be employed in the alternative.

[0044] If the first and second high-frequency power whose power varies in time as shown in FIG.3 is employed, the high-frequency power is applied intermittently to the reaction gas. For example, the first and second high-frequency powers are applied to the reaction gas from a time T2 to a time T3, and the high-frequency power is not applied to the reaction gas from a time T1 to a time T2. In the present embodiment, a time period during which the high-frequency power is applied, i.e., T3-T2 is 0.5×10^{-3} sec. Then, a time period during which the high-frequency power is not applied, i.e., T2-T1 is also 0.5×10^{-3} sec.

[0045] If the high-frequency powers whose power varies in time intermittently are employed, there is the possibility that the alkyl groups can be contained in the film in the complete form. That is, if the high-frequency powers whose power varies in time intermittently are applied, dissociation of the alkyl groups, which are contained in the compound having the siloxane bond (Si-O-Si), due to the application of the high-frequency powers can be suppressed during the time intervals in which the high-frequency powers is not applied.

[0046] Therefore, the alkyl groups which is not dissociate and thus of complete form can be contained in the insulating film 204. Accordingly, if the high-frequency powers whose power varies in time intermittently are applied, it is expected to be able to lower the dielectric constant of the insulating film 204 compared with the case where the high-frequency powers whose power is constant in time are applied.

[0047] Under the conditions shown in Table 1 and Table 2, or in case where the high-frequency powers whose power varies in time intermittently are applied under the conditions shown in Table 1 and Table 2, the methyl alcohol may be added into the reaction gas. Since the methyl groups can be contained much more in the film by the methyl alcohol, much more Si-O bonds in the film can be reduced. Therefore, it is expected that the dielectric constant of the film can be further lowered.

[0048] In order to add the methyl alcohol into the reaction gas, a flow rate of the liquid methyl alcohol is controlled by the liquid mass flowmeter (not shown), and then the liquid methyl alcohol is introduced into the chamber 101. Alternatively, the methyl alcohol may be vaporized by heating and then introduced into the chamber 101 in a gas state. When the methyl alcohol is added into the reaction gas, the flow rate of the methyl alcohol is 50 sccm.

②the case where the $\text{Si(OR)}_n\text{H}_m$ compound is used

[0049] The film forming conditions in this case will be given in Table 3.

Table 3

Flow rate of $\text{Si(OR)}_n\text{H}_m$ compound	50 sccm
Flow rate of an oxidizing gas	see Table 4
Flow rate of an inert gas (Ar or He)	200 sccm
Temperature of a substrate 103	200 °C
Pressure in a chamber 101	0.8 Torr
Frequency of the high frequency power applied by the first high frequency power supply 107	13.56 MHz
Power of the high frequency power applied by the first high frequency power supply 107	50 to 300 W
Frequency of the high frequency power applied by the second high frequency power supply 109	400 KHz
Power of the high frequency power applied by the second high frequency power supply 109	10 to 400 W

[0050] As the $\text{Si(OR)}_n\text{H}_m$ compound set forth in Table 3, there are $\text{Si(OCH}_3)_3\text{H}$, $\text{Si(OC}_2\text{H}_5)_3\text{H}$, $\text{Si(OCH}_3)_4$, and $\text{Si(OC}_2\text{H}_5)_4$, which are already mentioned above. Also, it should be noted that the $\text{Si(OR)}_n\text{H}_m$ compounds are not limited these specific example, but may use such a compounds satisfying $n+m=4$ and $m \geq 0$, where n and m denotes integer. Any one of these compounds may be contained in the reaction gas. These compounds, which are liquid at the room temperature (20°C), are heated to be vaporized after the flow rates of them are controlled by the liquid mass flowmeter (not shown), and then introduced into the chamber 101. Alternatively, the flow rates of these compounds may be controlled by the high-temperature mass flowmeter (not shown) after these compounds are vaporized, and then these compounds may be supplied to the chamber 101. The flow rate of the $\text{Si(OR)}_n\text{H}_m$ compound in Table 3 is the flow rate of the compound after vaporized.

[0051] Also, there are N_2O , O_2 , H_2O , CO_2 as the oxidizing gas set forth in the second line of Table 3, and at least one of them may be contained in the reaction gas. In case only one of these oxidizing gases is added into the reaction gas, i.e., without combination of other oxidizing gases, the flow rate of the oxidizing gas will be given in Table 4.

Table 4

Oxidizing gas	Flow rate
N_2O	50 to 200 scan
O_2	10 to 50 sccm
H_2O	100 to 300 scan
CO_2	100 to 300 scan

[0052] The dielectric constant of the insulating film 204 formed in accordance with these conditions was 2.5 at 1 MHz. This value is smaller than the dielectric constant 4.1 of the SiO_2 film used in the prior art. Also, the film forming rate was 200 nm/min.

[0053] The reason why the insulating film 204 having such a low dielectric constant can be formed is considered as follows. That is, if the $Si(OR)_nH_m$ compound is used, the alkoxy groups (OR) in the compound are contained in the film. In the insulating film 204, the Si-O bonds are disconnected at the portions where the alkoxy groups are contained, which in turn reduce the dielectric constant of that portions. As a result, the dielectric constant of the overall insulating film 204 is lowered.

[0054] In addition, Si-OR bonds have already been formed in the $Si(OR)_nH_m$ compound. Therefore, since Si and R (alkyl group) are easily bonded via O also in the insulating film 204, it can be prevented that Si and C are newly bonded to form the Si-C bonds. Accordingly, it is expected that, in the insulating film 204, the portions in which many Si-C bonds are continuously exist can be reduced compared with the case where the compound that does not have the Si-OR bond is employed. In this manner, the leakage current of the film can be reduced.

[0055] Under the conditions shown in Table 3 and Table 4, a C_pH_q gas may be further added into the reaction gas. If the C_pH_q gas is added, the methyl group, the ethyl group, etc. are contained in the film and thus the Si-O bonds in the film can be reduced. Therefore, it is expected that the dielectric constant of the film can be further lowered.

[0056] There are CH_4 , C_2H_4 , and C_2H_6 that are used actually as the C_pH_q gas in the experiment by the inventors of this application. However, the C_pH_q gas is not limited to these specific examples. At least one of these C_pH_q gases may be added to the reaction gas. In case only one of these gases is added into the reaction gas, i.e., without combination of other C_pH_q gases, the flow rates of these gases are 50 sccm.

(iii) Explanation of the plasma process performed to improve the hygroscopicity resistance of the insulating film

[0057] By the way, as an insulating film employed in semiconductor devices, it is preferable to use insulating films that show good hygroscopicity resistance. This is because if the insulating film that shows the poor hygroscopicity resistance and is apt to absorb the moisture is employed, the moisture in the film diffuses into the underlying wiring layer and thus corrosion of the wiring layer is caused.

[0058] Considering the above, in the film forming method of the present invention, in order to improve the hygroscopicity resistance of the insulating film 204 (see FIG.2B), the plasma process is performed for the insulating film 204 after the film is formed.

[0059] The conditions of this plasma process are given in Table 5 hereunder.

Table 5

(A)	(B)	(C)	(D)	(E)	(F)	(G)	(H)	(I)
O_2	600	--	--	400	50	400	0.2	60
N_2O	--	600	--	400	50	400	0.2	60

(B): O_2 flow rate (scan)

(C): N_2O flow rate (sccm)

(C): N_2O flow rate (sccm)

(E): Power of the first high-frequency power supply 107 (W)

(F): Power of the second high-frequency power supply 109 (W)

(G): Temperature of the substrate 103 ($^{\circ}C$)

(H): Pressure in the chamber 101 (Torr)

(I): Process time (sec)

Table 5 (continued)

(A)	(B)	(C)	(D)	(E)	(F)	(G)	(H)	(I)
NH ₃	--	--	600	400	50	400	0.2	60
O ₂ +N ₂ O	300	300	--	400	50	400	0.2	60
O ₂ +NH ₃	300	--	300	400	50	400	0.2	60
N ₂ O+NH ₃	--	300	300	400	50	400	0.2	60
O ₂ +N ₂ O+NH ₃	200	200	200	400	50	400	0.2	60

(A): Gas used in the plasma process

(B): O₂ flow rate (scan)(C): N₂O flow rate (sccm)

(E): Power of the first high-frequency power supply 107 (W)

(F): Power of the second high-frequency power supply 109 (W)

(G): Temperature of the substrate 103 (°C)

(H): Pressure in the chamber 101 (Torr)

(I): Process time (sec)

[0060] In this plasma process, a frequency of the high-frequency power applied by the first high-frequency power supply 107 is 13.56 MHz and a frequency of the high-frequency power applied by the second high-frequency power supply 109 is 400 kHz. As shown in the Table 5, there are O₂, N₂O, NH₃ as the gas employed in this plasma process, and at least one of them may be employed. In addition, Ar (argon) may be added to these gases. When Ar is added, the flow rate of Ar is 100 sccm.

[0061] If the plasma process is performed in this manner, H₂O that is contained in the insulating film 204 and CO₂ formed in this plasma process by oxidizing C in the film are discharged to the outside of the film. Also, since unbonded bonds of Si in the film are terminated by O, N, H, etc., the unbonded bonds of Si can be prevented from being bonded by OH group and the like, which in turn improve the hygroscopicity resistance of the film.

(iv) Explanation of an underlying insulating film and a cover insulating film formed to improve the hygroscopicity resistance of the insulating film

[0062] In order to improve the hygroscopicity resistance of the insulating film of the present invention, the underlying insulating film and the cover insulating film may be formed under and on the insulating film 204. This will be explained with reference to FIG.4. In FIG.4, the same reference symbols as those used in FIG.2 are affixed to the constituent members similar to those in FIG.2, and their explanation will be omitted hereunder.

[0063] In FIG.4, 205 denotes an underlying insulating film, and 206 denotes a cover insulating film. These films 205 and 206 are, for example, made up of SiO₂ and the like. In case where the SiO₂ film is employed as the underlying insulating film 205 and the cover insulating film 206, the film forming conditions of the SiO₂ film are given in Table 6 as follows.

Table 6

SiH ₄ flow rate	50 sccm
N ₂ O flow rate	1000 sccm
Temperature of the substrate 103	400 °C
Pressure in the chamber 101	0.8 Torr
Frequency of the high-frequency power applied by the first high frequency power supply 107	13.56 MHz
Power applied by the first high frequency power supply 107	50 W
Frequency of the high-frequency power applied by the second high frequency power supply 109	380 KHz
Power applied by the second high frequency power supply 109	400 W
Film thickness	50 nm

[0064] If the underlying insulating film 205 is formed in this manner, the moisture can be prevented from entering into the inside of the substrate 103 from the upper surface (the surface contacting to the underlying insulating film 205)

of the substrate 103. Therefore, corrosion of the aluminum wiring 203 by the moisture can be prevented.

[0065] Also, if the cover insulating film 206 is formed, the moisture can be prevented from entering into the inside of the insulating film 204 from the upper surface (the surface contacting to the cover insulating film 206) of the insulating film 204. Therefore, the degradation of the insulating film 204 due to the moisture can be prevented.

(v) Explanation of applying the present invention to the damascene process

[0066] In the above, the insulating film 204 of the present embodiment is formed on the aluminum wiring 203 (see FIG.2B). However, the present invention is not limited to this, and the insulating film 204 may be applied to the damascene process suitable for forming the copper wirings. Such damascene process will be explained with reference to FIGS.19A to 19F hereunder. FIGS.19A to 19F are sectional views showing the case where the insulating film 204 of the present invention is applied to the damascene process.

[0067] At first, as shown in FIG.19A, a lower copper wiring layer 210 is formed on an underlying interlayer insulating film 209, and a blocking film 211 such as the SiN film, etc. is formed on the lower copper wiring layer 210. The term blocking film is usually employed for the film when the film serves as both etching stopper film and barrier metal film.

[0068] Then, as shown in FIG.19B, the insulating film 204 of the present embodiment is formed on the blocking film 211. As previously explained, the insulating film 204 is formed by the plasma CVD method using ①the reaction gas containing the compound having the siloxane bond or ②the reaction gas containing the $\text{Si}(\text{OR})_n\text{H}_m$ compound. The conditions in this case are the same as those set forth in Table 1 to Table 4, which have already been described. The film thickness of the insulating film 204 can be controlled desirably by changing a film forming time.

[0069] Then, as shown in FIG.19C, a protection film 212 is formed on the insulating film 204. A dense and high quality silicon oxide film is used as the protection film 212. This protection film 212 serves to prevent the insulating film 204 from being deteriorated in an ashing process and etching process.

[0070] In turn, as shown in FIG.19D, photoresist 213 is coated on the protection film 212. After this, an opening portion 213a is formed in the photoresist 213 by the photolithography. Then, the protection film 212 and the insulating film 204 are etched by the RIE (Reactive Ion Etching) while using the photoresist 213 as a mask. In this RIE, the plasmanized gas mixture containing CF_4+CHF_3 or C_4F_8 may be used as the etching gas.

[0071] Then, the insulating film 204 located under the opening portion 213a in the photoresist 213 is removed by this etching, and an opening portion 204a is formed. Then, a surface of the blocking film 211 is exposed on the bottom portion of the opening portion 204a. Since the blocking film 211 has the etching resistance against the gas mixture containing CF_4+CHF_3 , the blocking film 211 is not etched by this RIE.

[0072] Then, as shown in FIG.19E, the blocking film 211 is etched by the RIE while using the photoresist 213 as a mask. In this RIE, a gas mixture containing CF_4+CHF_3 , in which the gas composition ratio is changed than that used in etching the insulating film 204, is plasmanized. By this plasmanised gas mixture, an opening portion 211a is formed in the blocking film 211 located under the opening portion 213a, and a surface of the lower copper wiring layer 210 is exposed. As a result, a via hole 214 reaching the lower copper wiring layer 210 is formed by this step.

[0073] Then, as shown in FIG.19F, after the photoresist 213 is removed and a barrier metal layer (not shown) such as a TiN film, etc. is formed on a side wall of the via hole 214, a copper plating film 215 is formed on an inner surface of the via hole 214 and an upper surface of the protection film 212. This copper plating film 215 is formed by supplying a current to a copper seed layer (not shown), which is formed on the inner surface of the via hole 214 and the upper surface of the protection film 212 by means of sputter method, etc.

[0074] After this step, the copper plating film 215 formed on the protection film 212 is removed by the CMP (Chemical Mechanical Polishing) method, and an upper copper wiring layer (not shown) is formed on the protection film 212.

[0075] According to the above mentioned damascene process, the insulating film 204 is formed as the interlayer insulating film between the lower copper wiring layer 210 and the upper copper wiring layer (not shown). This insulating film 204 is the low dielectric constant film that has the relative dielectric constant such as about 2.7 at 1 MHz, as described above. Therefore, if the insulating film 204 is employed, the wiring capacitance between the lower copper wiring layer 210 and the upper copper wiring layer (not shown) can be reduced, and therefore the semiconductor device that has the quick operation speed can be provided.

EXAMPLES

[0076] The inventors of this application examined how the property of the insulating film 204 changed when the film forming conditions were changed, especially when using HMDS (chemical formula: $\text{Si}(\text{CH}_3)_3\text{O}$) as the compound having the siloxane bond. The results of the examination will be explained in the following.

[0077] In this examination, a 8-inch silicon wafer was employed as the substrate 103 (see FIG.1). When the relative dielectric constant of the insulating film was measured, the C-V (Capacitance - Voltage) measuring method in which the high-frequency signal of 1 MHz was superposed on the DC bias was employed. When the refractive index of the

film was measured, the ellipsometer using the He-Ne laser whose wavelength is 632.8 nm was employed.

(1) The case where an electrode distance is 25 mm

5 [0078] At first, a distance (electrode distance) between the upper electrode 104 and the lower electrode 102 was fixed to 25 mm, and the film forming conditions were changed as follows.

(i) The case where the pressure was changed

10 [0079] The film forming conditions in this case are given as following Condition A.

(Condition A)

[0080]

15

HMDS flow rate: 50 sccm

N₂O flow rate: 200 sccm

substrate temperature: 350 °C

deposited film thickness: 500 nm

20

frequency of the high-frequency power applied by

the first high-frequency power supply 107: 13.56 MHz

power of the first high-frequency power supply 107: 300 W

power of the second power supply 109: 0 W

25

[0081] In the Condition A, the high-frequency power was applied only to the upper electrode 104 and was not applied to the lower electrode 102.

[0082] When the pressure of the atmosphere was changed under the Condition A, a relationship between the relative dielectric constant of the insulating film 204 and the pressure, and a relationship between the refractive index of the film and the pressure were obtained as in FIG.5. In FIG.5, an arrow pointing to the left indicates that the data series dotted by ● represent the relative dielectric constant, and an arrow pointing to the right indicates that the data series dotted by ○ represent the refractive index.

30

[0083] As can be seen from FIG.5, when the high-frequency power was applied only to the upper electrode 104, the relative dielectric constant was almost constant and its value showed about 2.8 even if the pressure of the atmosphere was changed. Similarly, the refractive index was almost constant and was about 1.38 to 1.42.

35

[0084] The leakage current of the insulating film 204 formed under the Condition A was of the order of 10^{-10} A/cm² at the electric field strength of 1 MV/cm over all pressures. This value is quite satisfactorily for practical use.

[0085] Next, the inventors of this application examined how the above tendency changed when the high-frequency power was applied not only to the upper electrode 104 but also to the lower electrode 102. The film forming conditions in this case are given as following Condition B.

40

(Condition B)

[0086]

45

HMDS flow rate: 50 sccm

N₂O flow rate: 200 sccm

substrate temperature: 350 °C

deposited film thickness: 500 nm

frequency of the high-frequency power applied by

50

the first high-frequency power supply 107: 13.56 MHz

power of the first high-frequency power supply 107: 300 W

frequency of the high-frequency power applied by

the second high-frequency power supply 109: 380 KHz

power of the second power supply 109: 30 W

55

[0087] When the pressure of the atmosphere was changed under the Condition B, a relationship between the relative dielectric constant of the insulating film 204 and the pressure, and a relationship between the refractive index of the film and the pressure were obtained as in FIG.6.

[0088] As can be seen from FIG.6, when the high-frequency power was applied to both the upper electrode 104 and the lower electrode 102, the tendency that is clearly different from that shown in FIG.5 was obtained. That is, when the high-frequency power was applied to both electrodes, such a tendency appeared that, when the pressure was increased, the relative dielectric constant and the refractive index of the film were reduced.

5 [0089] In FIG.6, the value of the relative dielectric constant at the pressure of 1.0 Torr was about 3.8. In the present example, the measurement was not carried out at the pressure lower than 1.0 Torr. However, it can be understood from FIG.6 such a tendency that the relative dielectric constant comes close to 4.0 at the pressure lower than 1.0 Torr.

[0090] As described in the prior art column, in order to provide the semiconductor device in which the wiring capacitance is reduced smaller than the prior art, the film whose relative dielectric constant is lower than that of SiO₂ film (4.1) must be employed as the interlayer insulating film. It can be understood from the above description that, to this end, the pressure should be set to more than 1.0 Torr and the high-frequency power should be applied to both the upper electrode 104 and the lower electrode 102.

[0091] Though the electrode distance was set to 25 mm in this example, the inventors of this application similarly examined the case where the electrode distance was set to more than 50 mm. According to this examination, it became apparent that, when the electrode distance was set to more than 50 mm, the insulating film 204 having the relative dielectric constant lower than that of the SiO₂ film (4.1) could be formed by setting the pressure to more than 0.5 Torr.

[0092] The leakage current of the insulating film 204 formed under the Condition B was of the order of 10⁻¹⁰ A/cm² at the electric field strength of 1 MV/cm over all pressures. This value is quite satisfactorily for practical use.

20 (ii) The case where the power of the second high-frequency power supply 109 was changed

[0093] As described above, it was found that, when the high-frequency power was applied not only to the upper electrode 104 but also to the lower electrode 102, the tendency that was different from the case where the high-frequency power is applied only to one electrode was obtained.

25 [0094] Next, the inventors of this application examined what change occurred on the property of the insulating film 204 when the power of the high-frequency power applied to one electrode was changed while applying the high-frequency power to the other electrode.

[0095] The film forming conditions in this case are given as following Condition C.

30 (Condition C)

[0096]

35 HMDS flow rate: 50 sccm
N₂O flow rate: 200 sccm
substrate temperature: 350 °C
deposited film thickness: 500 nm
frequency of the high-frequency power applied by
the first high-frequency power supply 107: 13.56 MHz
40 power of the first high-frequency power supply 107: 100 W
frequency of the high-frequency power applied by
the second high-frequency power supply 109: 380 KHz

45 [0097] When the power of the second high-frequency power was changed under the Condition C, a relationship between the power of the second high-frequency power supply 109 and a dynamic hardness of the insulating film 204, and a relationship between the power of the second high-frequency power supply 109 and a Young's modulus of the insulating film 204 were obtained as shown in FIG.7. The measurement of the dynamic hardness and Yang's modulus is carried out using the equipment DUH-W201S, which is made by Shimadzu Corporation.

50 [0098] As can be clearly seen from FIG.7, the dynamic hardness and the Young's modulus of the film increases as the power of the second high-frequency power supply 109 increases. Put in another way, when the power of the second high-frequency power supply 109 is larger than 0 W, the film whose dynamic hardness and Young's modulus are larger than that obtained when the power is 0 W can be obtained. This means that if the high-frequency power is applied not only to the upper electrode 104 but also to the lower electrode 102, the film whose dynamic hardness and Young's modulus are larger than that obtained when the power is applied only to the upper electrode 104 can be obtained.

55 [0099] By the way, in the damascene process for forming the copper wiring layer, the damascene trench is formed in the interlayer insulating film, and the copper plating is preformed for the upper surface of the interlayer insulating film and the inside of the damascene trench. After this, in order to leave the copper only in the damascene trench, overall surface of the copper plating is polished by the CMP method. When polishing the surface by the CMP method,

the interlayer insulating film, together with the copper plating thereon, is polished to some extent. If the interlayer insulating film is polished in this manner, the residual film thickness of the interlayer insulating film must be desirably controlled. In general, it has been known that in order to control the residual film thickness in the CMP, the film having the large dynamic hardness is preferable for the interlayer insulating film

[0100] As described above, if the high-frequency power is applied to both the upper electrode 104 and the lower electrode 102, the film whose dynamic hardness is larger than that obtained when the high-frequency power is applied only to the upper electrode 104 can be formed. Accordingly, it is preferable to use the insulating film 204, formed by applying the high-frequency power to both the upper and lower electrode 104 and 102, for the insulating film which undergoes the polishing in the CMP process.

[0101] The leakage current of the insulating film 204 formed under the Condition C was of the order of 10^{-10} A/cm² at the electric field strength of 1 MV/cm over all pressures. This value is quite satisfactorily for the practical use.

(2) The case where the electrode distance is 50 mm

[0102] The results of the examination set forth in the case (1) were derived when the electrode distance was fixed to 25 mm and the film forming conditions (except the electrode distance) were changed. However, it is considered that the property of the insulating film 204 may also change as the electrode distance changes. Therefore, the inventors of this application examined how the property of the insulating film 204 changed by changing the electrode distance. In particular, the examination was carried out while fixing the electrode distance to 50 mm in the following.

(i) The case where the high-frequency power is applied only to the upper electrode 104 and the power of the high-frequency power is changed

[0103] The film forming conditions in this case are given as following Condition D.

(Condition D)

[0104]

HMDS flow rate: 50 sccm
 N₂O flow rate: 200 sccm
 substrate temperature: 350 °C
 deposited film thickness: 500 nm
 pressure: 0.9 Torr
 frequency of the high-frequency power applied by
 the first high-frequency power supply 107: 13.56 MHz
 power of the second high-frequency power supply 109: 0 W

[0105] As can be understood from the Condition D, the high-frequency power was applied only to the upper electrode 104, and the high-frequency power was not applied to the lower electrode 102.

[0106] When the power of the first high-frequency power supply 107 was changed under the Condition D, a relationship between the power of the first high-frequency power supply 107 and the relative dielectric constant of the insulating film 204, and a relationship between the power and the refractive index of the insulating film 204 were obtained as shown in FIG.8.

[0107] As can be seen from FIG.8, it can be understood that if the power of the high-frequency power was changed, the relative dielectric constant and the refractive index of the insulating film 204 were seldom changed. More specifically, the relative dielectric constant of the insulating film 204 is about 2.6 and the refractive index is about 1.38.

[0108] The leakage current of the insulating film 204 formed under the Condition D was of the order of 10^{-10} A/cm² at the electric field strength of 1 MV/cm over all the powers of the first high-frequency power supply 107. This value is quite satisfactorily for the practical use.

(ii) The case where the high-frequency power is applied to both the upper electrode 104 and the lower electrode 102, and the power of the high-frequency power applied to the lower electrode 102 is changed

[0109] The film forming conditions in this case are given as following Condition E.

(Condition E)

[0110]

5 HMDS flow rate: 50 sccm
 N₂O flow rate: 200 sccm
 substrate temperature: 350 °C
 deposited film thickness: 500 nm
 pressure: 0.9 Torr
 10 frequency of the high-frequency power applied by
 the first high-frequency power supply 107: 13.56 MHz
 power of the first high-frequency power supply 107: 300 W
 frequency of the high-frequency power applied by
 the second high-frequency power supply 109: 380 KHz
 15

[0111] When the power of the second high-frequency power supply 109 was changed under the Condition E, a relationship between the power of the second high-frequency power supply 109 and the relative dielectric constant of the insulating film 204, and a relationship between the power and the refractive index of the insulating film 204 were obtained as shown in FIG.9.

20 [0112] As can be seen from FIG.9, it can be understood that, if the power of the second high-frequency power supply 109 is increased, the relative dielectric constant and the refractive index of the film are increased correspondingly. However, it can be understood from FIG.9 that the relative dielectric constant and the refractive index have their maximum values when the power of the second high-frequency power supply 109 is around 100 W, and they are reduced when the power is larger than 100 W. Also, it can be seen from FIG.9 that the relative dielectric constant does not exceed 4.0 at the maximum value. Accordingly, even if the high-frequency power is applied to the upper electrode 104 and the lower electrode 102, the relative dielectric constant of the insulating film 204 does not exceed the relative dielectric constant (4.1) of the conventional SiO₂ film.

25 [0113] Here, in order to see how the relative dielectric constant of the insulating film 204 changed when the electrode distance was changed, FIG.5 and FIG.9 are compared in the following. Particularly, in order to avoid disparity in the condition other than the electrode distance, the relative dielectric constant (about 2.9) obtained when the pressure is 1.0 Torr in FIG.5 should be compared with the relative dielectric constant (about 2.6) obtained when the power of the second high-frequency power supply 109 is 0 W in FIG.9. These values were obtained respectively when the film forming conditions other than the electrode distance were set substantially equal. More specifically, these values were obtained when only the power of the first high-frequency power supply 107, which was 300 W, was applied and the pressure was about 1.0 Torr.

30 [0114] As can be seen clearly by comparing these values, the dielectric constant obtained when the electrode distance was set to 50 mm became lower than that obtained when the electrode distance was set to 25 mm. The reason for this will be given as follows. That is, if the electrode distance is large, the area (sheath area) where the gradient of the potential appears between the electrodes is relatively reduced compared with the case where the electrode distance is narrow. If the sheath area is reduced in this manner, the acceleration of the methyl groups caused by this sheath area can be suppressed and decomposition of the methyl groups can also be suppressed. Therefore, the methyl groups can be taken into the insulating film 204 with their complete form, and therefore the dielectric constant of the insulating film 204 is lowered.

35 [0115] Therefore, in order to lower the dielectric constant of the insulating film 204, it is preferable to expand the electrode distance. More specifically, it is preferable to set the electrode distance more than 25 mm.

40 [0116] The leakage current of the insulating film 204 formed under the Condition E was of the order of 10⁻¹⁰ A/cm² at the electric field strength of 1 MV/cm at all powers of the second high-frequency power supply 109. This value is quite satisfactorily for practical use.

45 (3) The case where H₂O is employed as an oxidizing gas

50 [0117] The above results of the examination were obtained by employing N₂O as the oxidizing gas. The inventors of this application examined how the property of the insulating film 204 changed when the film forming conditions were changed when using H₂O as the oxidizing gas in place of N₂O.

55 (i) The case where the high-frequency power is applied only to the lower electrode 102 and the pressure is changed

[0118] The film forming conditions in this case are given as following Condition F.

(Condition F)

[0119]

5 HMDS flow rate: 50 sccm
 H₂O flow rate: 100 sccm
 substrate temperature: 200 °C
 deposited film thickness: 500 nm
 electrode distance: 25 mm
 10 power of the first high-frequency power supply 107: 0 W
 frequency of the high-frequency power applied by the second high-frequency power supply 109: 380 KHz power
 of the second high-frequency power supply 109: 100 W

15 [0120] As can be understood from the Condition F, the high-frequency power was applied only to the lower electrode 102 and was not applied to the upper electrode 104.

[0121] When the pressure was changed under the Condition F, a relationship between the relative dielectric constant of the insulating film 204 and the pressure, and a relationship between the refractive index and the pressure were obtained as shown in FIG.10.

20 [0122] As can be seen from FIG.10, it can be understood that the relative dielectric constant and the refractive index of the insulating film 204 are monotonously reduced as the pressure of the atmosphere is increased. It can also be understood that the relative dielectric constant of the insulating film 204 is lower than that of the conventional SiO₂ film (4.1) at all pressures.

[0123] The leakage current of the insulating film 204 formed under the Condition F was of the order of 10⁻¹⁰ A/cm² at the electric field strength of 1 MV/cm at all pressures. This value is quite satisfactorily for practical use.

25 (ii) The examination results of H₂O contained in the film

[0124] If H₂O is used as the oxidizing gas as above, there is the possibility that H₂O is contained in the insulating film 204. The inventors of this application therefore examined how the H₂O contained in the insulating film 204 changed when the pressure was changed under the Condition F.

[0125] The results of this examination are shown in FIG.11. FIG.11 shows the measurement results measured by the Infrared Absorption Spectroscopy.

35 [0126] If H₂O is contained in the film, a peak appears in vicinity of the wave number of 3600 cm⁻¹. However, as can be seen from FIG.11, no peak appears in vicinity of 3600 cm⁻¹ at all pressures. This means that regardless of the pressure of the atmosphere, no H₂O is contained in the insulating film 204 that is formed in accordance with the Condition F.

[0127] In general, if H₂O is contained in the interlayer insulating film, H₂O diffuses into the underlying wiring layer formed under the interlayer insulating film, and such a problem arises that the wiring layer corrodes due to the diffusion of the H₂O. As described above, in the present embodiment, even if H₂O is used as the oxidizing gas, no H₂O is contained in the insulating film 204, and thus such problem does not arise.

40 [0128] In addition, according to the results of another examination made by the inventors of this application, such a good result was obtained that the film thickness uniformity of the insulating film 204 is less than 3 %.

45 (iii) The case where the high-frequency power is applied also to the upper electrode 104, and the power of the high-frequency power applied to the lower electrode 102 is changed

[0129] In the Condition F, the high-frequency power is applied only to the lower electrode 102. The inventors of this application examined how the property of the insulating film 204 changed when the high-frequency power was applied also to the upper electrode 104 and the power of the high-frequency power applied to the lower electrode 102 was changed.

[0130] The film forming conditions in this case are given as following Condition G.

(Condition G)

55 [0131]

HMDS flow rate: 50 sccm
 H₂O flow rate: 250 sccm

substrate temperature: 375 °C
 deposited film thickness: 500 nm
 electrode distance: 25 mm
 pressure: 2.3 Torr

frequency of the high-frequency power applied by
 the first high-frequency power supply 107: 13.56 MHz
 power of the first high-frequency power supply 107: 300 W
 frequency of the high-frequency power applied by
 the second high-frequency power supply 109: 380 KHz

[0132] When the relative dielectric constant and the refractive index of the resultant insulating film 204 examined while changing the power of the high-frequency power applied by the second high-frequency power supply 109 under Condition G, the results shown in FIG.12 were obtained.

[0133] As shown in FIG.12, it can be understood that as the power of the second high-frequency power supply 109 is reduced, the relative dielectric constant and the refractive index of the insulating film 204 are lowered.

[0134] Next, the inventors of this application examined the leakage current of the insulating film 204 formed under the Condition G by changing variously the power of the second high-frequency power supply 109 as in the above.

[0135] In this examination, as shown in FIG.13, the insulating film 204 was formed on a p-type silicon substrate 207 under the Condition G. Then, with the p-type silicon substrate 207 being grounded, a mercury probe 208, which has an electrode area of 0.02258 cm², was brought into contact with the upper surface of the insulating film 204, and a negative potential is given to the mercury probe 208.

[0136] Results of the examination are shown in FIG.14. In FIG.14, an ordinate represents the leakage current (A/cm²) of the insulating film 204 in a logarithmic scale. On the other hand, an abscissa represents the strength of the electric field (MV/cm) applied to the insulating film 204, and a (-) sign indicates that the negative potential is applied to the mercury probe 208.

[0137] As shown in FIG.14, the leakage current is 10⁻¹⁰ to 10⁻⁹ A/cm² at -1 MV/cm, and this value is quite satisfactorily for practical use.

(iv) Examination results of NH₃ contained in the film

[0138] It is preferable that, when the insulating film 204 is used as the interlayer insulating film and chemical amplification resist is coated on the insulating film 204 for patterning, NH₃ is not contained in the insulating film 204. This is because, if NH₃ is contained in the insulating film 204, the chemical amplification resist formed on the insulating film 204 is crosslinked by the NH₃ when the resist undergoes patterning, and thus the desired resist pattern cannot be formed. Especially, in case where the fine pattern is required for the resist, this phenomenon becomes prominent, which poses a barrier to implement fine patterning for the underlying insulating film.

[0139] When the nitrogen oxide (N₂O, etc.), which serves as the oxidizing gas, is added into the reaction gas, NH₃ is apt to be contained in the film because of N (nitrogen) in the nitrogen oxide. In order to verify this phenomenon, the inventors of this application formed the insulating film by using Si(CH₃)₄ and N₂O, and then examined an amount of NH₃ in the insulating film. The film forming conditions for this insulating film are given as following Condition H.

(Condition H)

[0140]

Si(CH₃)₄ flow rate: 80 sccm
 N₂O flow rate: 320 sccm
 substrate temperature: 350 °C
 deposited film thickness: 500 nm
 pressure: 1 Torr

frequency of the high-frequency power applied by
 the first high-frequency power supply 107: 13.56 MHz
 power of the first high-frequency power supply 107: 300 W
 frequency of the high-frequency power applied by the second high-frequency power supply 109: 400 KHz power
 of the second high-frequency power supply 109: 30 W

[0141] The results of an amount of NH₃ contained in the insulating film formed under the Condition H are shown in FIG.15. FIG.15 is graph showing the results of an amount of NH₃ measured by the TDS (Thermal Desorption Spec-

troscopy) method. An abscissa of this graph represents the temperature (°C) at which the insulating film is heated in the measurement. On the other hand, an ordinate represents the relative ion intensity (%) of monovalent positive ion whose molecular weight is 17. Here, the relative ion intensity (%) is defined as (ion intensity of the monovalent positive ion whose molecular weight is 17)/(total ion intensity desorbed by heating). NH_3 is ionized to NH_3^+ by the heating and since the NH_3^+ is monovalent positive ion whose molecular weight is 17, what depicted in the FIG.15 is the relative ion intensity of the NH_3^+ .

[0142] As can be seen from FIG.15, it can be understood that when the bias is applied (which means the power of the second high-frequency power supply 109 is applied to the lower electrode 102 as in the condition H), NH_3^+ is desorped at the temperature of about 250 °C. It can also be understood that when no bias is applied (which means the power of the second high-frequency power supply 109 is not applied to the lower electrode 102), NH_3^+ is desorped at the temperature of about 400 °C.

[0143] In this manner, it can be understood that if the nitrogen oxide (N_2O , etc.) is used as the oxidizing gas, NH_3 is contained in the film regardless whether the bias is applied or not.

[0144] On the contrary, it is considered that if H_2O is used as the oxidizing gas, NH_3 is not contained in the film. In order to verify this, the inventors of this application examined an amount of NH_3 contained in the insulating film 204 that is formed under the following Condition I.

(Condition I)

[0145]

HMDS flow rate: 50 sccm

H_2O flow rate: 250 sccm

substrate temperature: 375 °C

deposited film thickness: 500 nm

electrode distance: 25 mm

frequency of the high-frequency power applied by the first high-frequency power supply 107: 13.56 MHz power of the first high-frequency power supply 107: 300 W

power of the second high-frequency power supply 109: 0 W

[0146] As can be seen from the Condition I, the high-frequency power is applied only to the upper electrode 104 and is not applied to the lower electrode 102. That is, bias is not applied in the Condition I.

[0147] When the pressure in forming the film was changed variously under the Condition I, an amount of NH_3 contained in the insulating film 204 was measured as shown in FIG.16. FIG.16 is a graph showing the results obtained when the amount of NH_3 contained in the insulating film was measured by the TDS method. Since the abscissa and the ordinate of this graph are the same as those explained previously in FIG.15, their explanation will be omitted.

[0148] Although the desorption begins at the temperature of 600 °C in FIG.16, this is not due to NH_3^+ but due to the isotope of CH_4^+ whose molecular weight is 17. In this manner, if H_2O is used as the oxidizing gas, the amount of NH_3 contained in the film can be largely reduced compared with the case where N_2O is used as the oxidizing gas. Therefore, if the insulating film 204 is formed by using H_2O as the oxidizing gas and the chemical amplification resist is formed thereon, the chemical amplification resist can be patterned with desired precision. This respect will be explained with FIGS.20A to 20G, in which the damascene process is exemplified. FIGS.20A to 20G are sectional views showing a sectional shape respectively when the damascene process is applied to the insulating film 204 formed by using H_2O as the oxidizing gas.

[0149] First, as shown in FIG.20A, a lower copper wiring layer 210 is formed on an underlying interlayer insulating film 209 and then a blocking film 211 such as an SiN film, etc. is formed thereon.

[0150] Then, as shown in FIG.20B, the insulating film 204 is formed on the blocking film 211 under the Condition I. At this time, as described above, an amount of NH_3 contained in the insulating film 204 is sufficiently reduced.

[0151] Then, as shown in FIG.20C, chemical amplification resist 216 is coated on the insulating film 204.

[0152] Then, as shown in FIG.20D, an opening portion 216a is formed by patterning the chemical amplification resist 216 by virtue of the photolithography. As described above, since an amount of NH_3 contained in the insulating film 204 has been sufficiently reduced, the chemical amplification resist 216 is not crosslinked by NH_3 during undergoing this patterning. Therefore, fine pattern of the chemical amplification resist 216 can be obtained in this step, and thus a diameter of the opening portion 216a can be made small desirably.

[0153] Then, as shown in FIG.20E, using the chemical amplification resist 216 as an etching mask, an opening portion 204a is formed by etching the insulating film 204. This etching is carried out by the RIE, in which a gas mixture containing $\text{CF}_4 + \text{CHF}_3$ or C_4F_8 is used as the etching gas. Since the pattern of the chemical amplification resist 216 is formed finely as described above, the pattern of the insulating film 204 can also be formed finely. Thus, the opening

portion 204a with a small diameter can be formed in the insulating film 204. In this etching, the blocking film 211 has etching resistance against the gas mixture containing $\text{CF}_4 + \text{CHF}_3$, which is used as etching gas. Therefore, the blocking film 211 is not etched by this etching.

[0154] Then, as shown in FIG.20F, using the chemical amplification resist 216 as a etching mask, the blocking film 211 is etched by the RIE. In this RIE, a gas mixture containing $\text{CF}_4 + \text{CHF}_3$, in which the gas composition ratio is changed than that used in etching the insulating film 204, is plasmanized. By this plasmanised gas mixture, an opening portion 211a is formed in the blocking film 211 located under the opening portion 216a, and a surface of the lower copper wiring layer 210 is exposed. As a result, a via hole 214 reaching the lower copper wiring layer 210 is formed by this step.

[0155] Then, as shown in FIG.20G, after the chemical amplification resist 216 is removed, a barrier metal layer (not shown) such as the TiN film, etc. is formed on a side wall of the via hole 214, and then a copper plating film 215 is formed in the inside of the via hole 214 and on the insulating film 204. This copper plating film 215 is formed by supplying a current to a copper seed layer (not shown), which is formed on the inner surface of the via hole 214 and the upper surface of the insulating film 204 by means of sputter method, etc.

[0156] After this step, the copper plating film 215 formed on the insulating film 204 is removed by the CMP (Chemical Mechanical Polishing) method, and then an upper copper wiring layer (not shown) is formed on the insulating film 204.

[0157] As described above, if the insulating film 204 is formed by using the reaction gas containing HMDS and H_2O , and then the chemical amplification resist 216 is formed thereon, the chemical amplification resist 216 is not crosslinked by NH_3 . Therefore, since the chemical amplification resist 216 can be patterned finely, the insulating film 204 can be etched finely using the chemical amplification resist 216 as a etching mask.

[0158] In this fashion, if the insulating film 204 is formed by using the reaction gas containing HMDS and H_2O , not only the dielectric constant of the film 204 can be lowered, but also the amount of NH_3 in the film 204 can be reduced and therefore the fine pattern can easily be made in the film 204.

[0159] Next, the inventors of this application examined the relative dielectric constant and the refractive index of the insulating film 204 obtained when the pressure of the atmosphere was changed under the Condition I. The examination results are shown in FIG.17.

[0160] As shown in FIG. 17, it can be understood that as the pressure is increased, both the relative dielectric constant and the refractive index reduces.

[0161] In addition, the inventors of this application examined the leakage current in the insulating film 204 formed under the same conditions as FIG.17. Since the leakage current measuring method is the same as that explained with reference to FIG.13, its explanation will be omitted.

[0162] The examination results are shown in FIG. 18. As shown in FIG.18, the leakage current is 10^{-10} to 10^{-9} A/cm² at -1 MV/cm, and this value is quite satisfactorily 0 for practical use.

[0163] As described above, according to the film forming method of the present invention, the insulating film containing silicon is formed by using the compound having the siloxane bond or the $\text{Si}(\text{OR})_n\text{H}_m$ compound. The alkoxy or alkyl group is contained in the insulating film, and the dielectric constant of the portion of the film in which The alkoxy or alkyl group is contained is lowered, and thus the dielectric constant of the entire film is lowered.

[0164] Also, in the compound having the siloxane bond or the $\text{Si}(\text{OR})_n\text{H}_m$ compound, Si and O have already been bonded. Therefore, when the insulating film containing silicon is formed by using these compounds, many Si-C bonds are hard to be formed in the film. Therefore, the increase in the leakage current due to the Si-C bond can be reduced in the insulating film containing silicon formed as above.

[0165] Especially, when the parallel plate type plasma chemical vapor deposition equipment is used as the film forming equipment, and also the gas containing HMDS (chemical formula: $(\text{Si}(\text{CH}_3)_3)_2\text{O}$) and N_2O is used as the reaction gas, following particular advantages can be achieved respectively if the film is formed under following conditions.

(1) Applying the high-frequency power to both the upper and lower electrode.

[0166] According to this condition, the insulating film containing silicon having the larger dynamic hardness than the case where the high- frequency power is applied only to the upper electrode can be formed.

(2) Applying the high-frequency power to both the upper and lower electrode, and setting the pressure of the atmosphere to more than 0.5 Torr.

[0167] According to this condition, the insulating film containing silicon having the larger dynamic hardness than the case where the high- frequency power is applied only to the upper electrode and having the lower dielectric constant than that of the SiO_2 film can be formed.

(3) Applying the high-frequency power only to the upper electrode, and setting the electrode distance to more than 25 mm.

[0168] According to this condition, the insulating film containing silicon having the lower dielectric constant than the case where the high-frequency power is applied only to the upper electrode and the distance between the upper and lower electrodes is less than 25 mm can be formed.

[0169] Also, if the reaction gas containing HMDS and H₂O is used, NH₃ is seldom contained in the resultant insulating film containing silicon. Therefore, the chemical amplification resist formed on the insulating film containing silicon is never crosslinked by NH₃. As a result, the resist can be finely patterned and thus the insulating film containing silicon can be finely patterned desirably.

[0170] The invention is not to be constructed as limited to the particular examples describes herein, as these are to be regarded as illustrative, rather than restrictive. The invention is intended to cover all process and structures which do not depart from the spirit and scope of the invention. For example, though in the examples and examination therein the HMDS are utilized, other compounds having the siloxane bonds such as OMCTS, HEDS, TMDS, TEDS, TMCTS, TETCS may be utilizing alternatively.

Claims

1. A film forming method of forming an insulating film containing silicon on a substrate by plasmanizing a compound having siloxane bonds and an oxidizing gas to react with each other.
2. A film forming method according to claim 1, wherein the compound having the siloxane bonds is any one selected from the group consisting of (Si(CH₃)₃)₂O, (Si(CH₃)₂)₄O₄, (Si(C₂H₅)₃)₂O, (SiH(CH₃)₂)₂O, (SiH(C₂H₅)₂)₂O, (SiH(CH₃))₄O₄, and (SiH(C₂H₅))₄O₄.
3. A film forming method according to claim 2, wherein the oxidizing gas contains at least one selected from the group consisting of N₂O, O₂, H₂O, and CO₂.
4. A film forming method according to any one of claim 1 to claim 3, wherein an inert gas is further added in addition to the compound having the siloxane bonds and the oxidizing gas.
5. A film forming method according to any one of claim 1 to claim 4, wherein plasmanization is performed by applying a high-frequency power to the compound having the siloxane bonds and the oxidizing gas.
6. A film forming method according to claim 5, wherein the high-frequency power is applied intermittently to the compound having the siloxane bond and the oxidizing gas.
7. A film forming method of performing a film formation in a parallel plate type plasma chemical vapor deposition equipment in which an upper electrode and a lower electrode are provided to oppose to each other in a chamber, wherein an insulating film containing silicon is formed on a substrate in an atmosphere in which a high-frequency power is applied to the upper electrode and the lower electrode and a reaction gas containing (Si(CH₃)₃)₂O and N₂O is introduced into the chamber.
8. A film forming method according to claim 7, wherein a pressure of the atmosphere is set to more than 0.5 Torr.
9. A film forming method of performing a film formation in a parallel plate type plasma chemical vapor deposition equipment in which an upper electrode and a lower electrode are provided to oppose to each other in a chamber, wherein an insulating film containing silicon is formed on a substrate in an atmosphere in which a distance between the upper electrode and the lower electrode is set to more than 25 mm, a high-frequency power is applied only to the upper electrode, and a reaction gas containing (Si(CH₃)₃)₂O and N₂O is introduced into the chamber.
10. A film forming method of forming an insulating film containing silicon on a substrate by plasmanizing a Si(OR)_nH_m compound (where R is an alkyl group, n and m denotes integer satisfying n+m=4 and m≥0) and an oxidizing gas to react to each other.
11. A film forming method according to claim 10, wherein the Si(OR)_nH_m compound is any one selected from the group consisting of Si(OCH₃)₃H, Si(OC₂H₅)₃H, Si(OCH₃)₄, and Si(OC₂H₅)₄.

12. A film forming method according to claim 10 or claim 11, wherein the oxidizing gas contains at least one selected from the group consisting of N_2O , O_2 , H_2O , and CO_2 .
13. A film forming method according to any one of claim 10 to claim 12, wherein a C_pH_q compound is further added in addition to the $\text{Si}(\text{OR})_n\text{H}_m$ compound and the oxidizing gas.
14. A film forming method according to claim 13, wherein the C_pH_q compound is any one selected from the group consisting of CH_4 , C_2H_4 , and C_2H_6 .
15. A film forming method according to any one of claim 10 to claim 14, wherein an inert gas is further added in addition to the $\text{Si}(\text{OR})_n\text{H}_m$ compound and the oxidizing gas.
16. A film forming method according to any one of claim 1 to claim 15, wherein an atmosphere containing at least one of O_2 , N_2O , and NH_3 is plasmanized after the insulating film containing silicon is formed, and then a surface of the insulating film containing silicon is exposed to the plasmanized atmosphere.
17. A film forming method according to any one of claim 1 to claim 16, wherein an underlying insulating film is formed on the substrate and the insulating film containing silicon is formed on the underlying insulating film.
18. A film forming method according to any one of claim 1 to claim 17, wherein a cover insulating film is formed after the insulating film containing silicon is formed.
19. A semiconductor device comprising the insulating film containing silicon formed by employing the film forming method set forth in claim 1.

FIG. 1

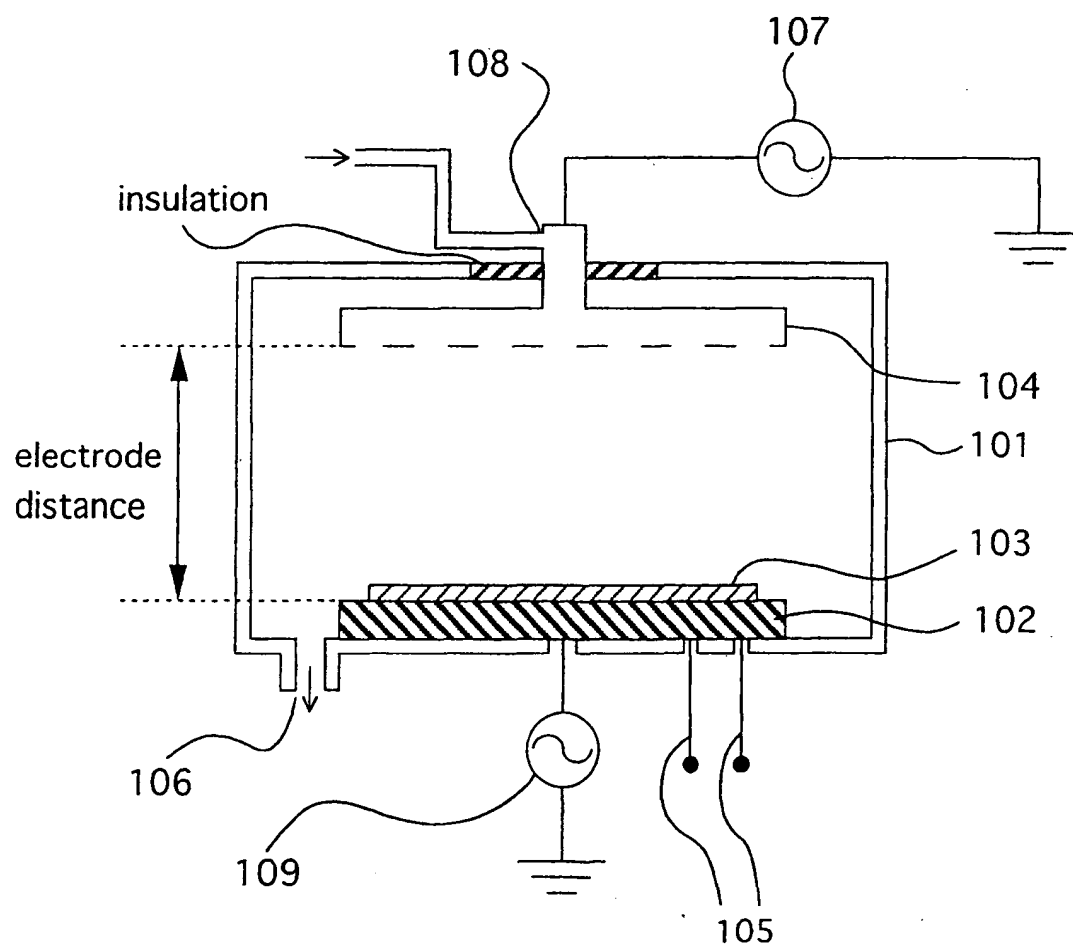


FIG.2A

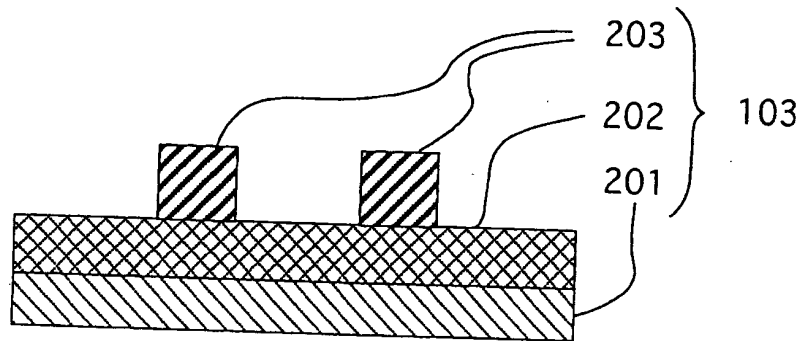


FIG.2B

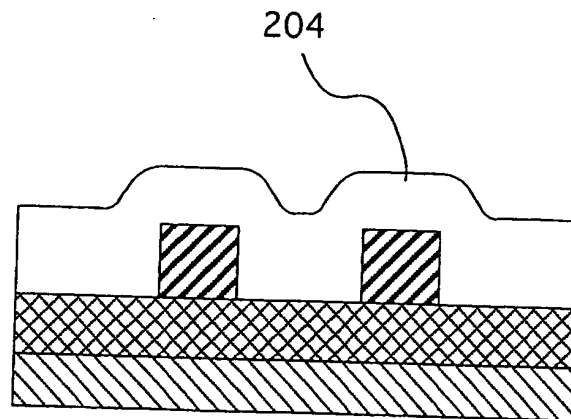


FIG.3

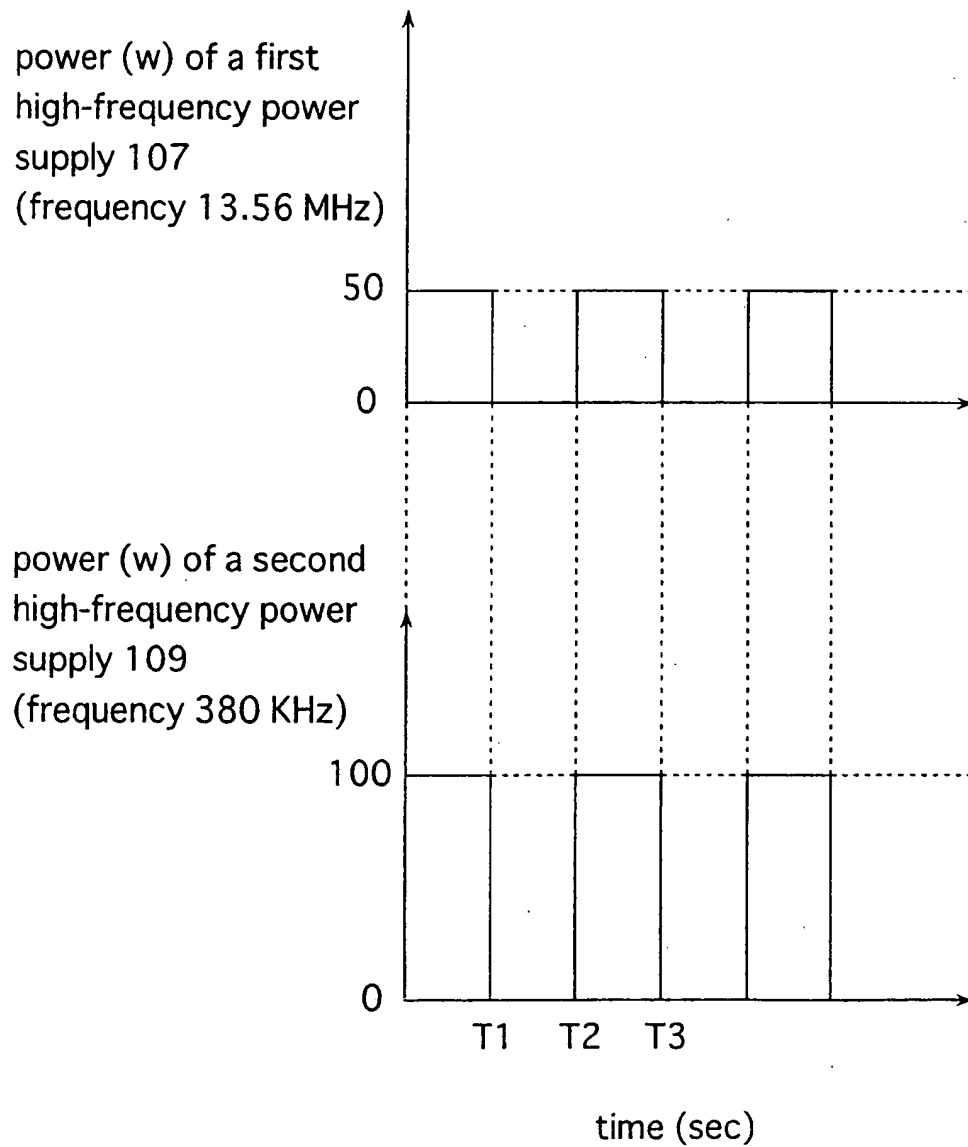


FIG.4

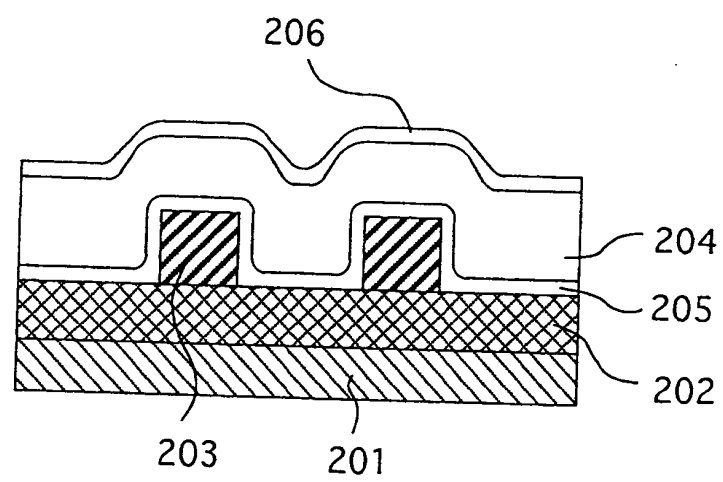
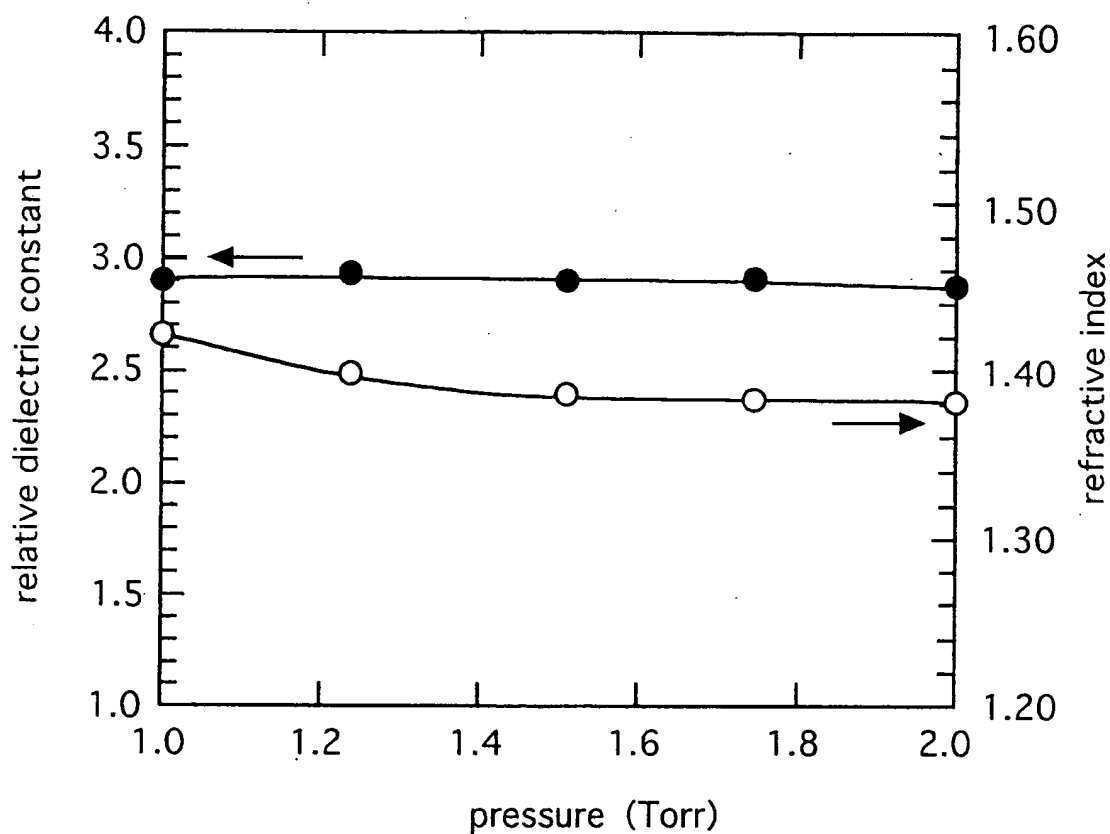


FIG.5



HMDS flow rate : 50sccm

N₂O flow rate : 200sccm

substrate temperature : 350°C

electrode distance : 25mm

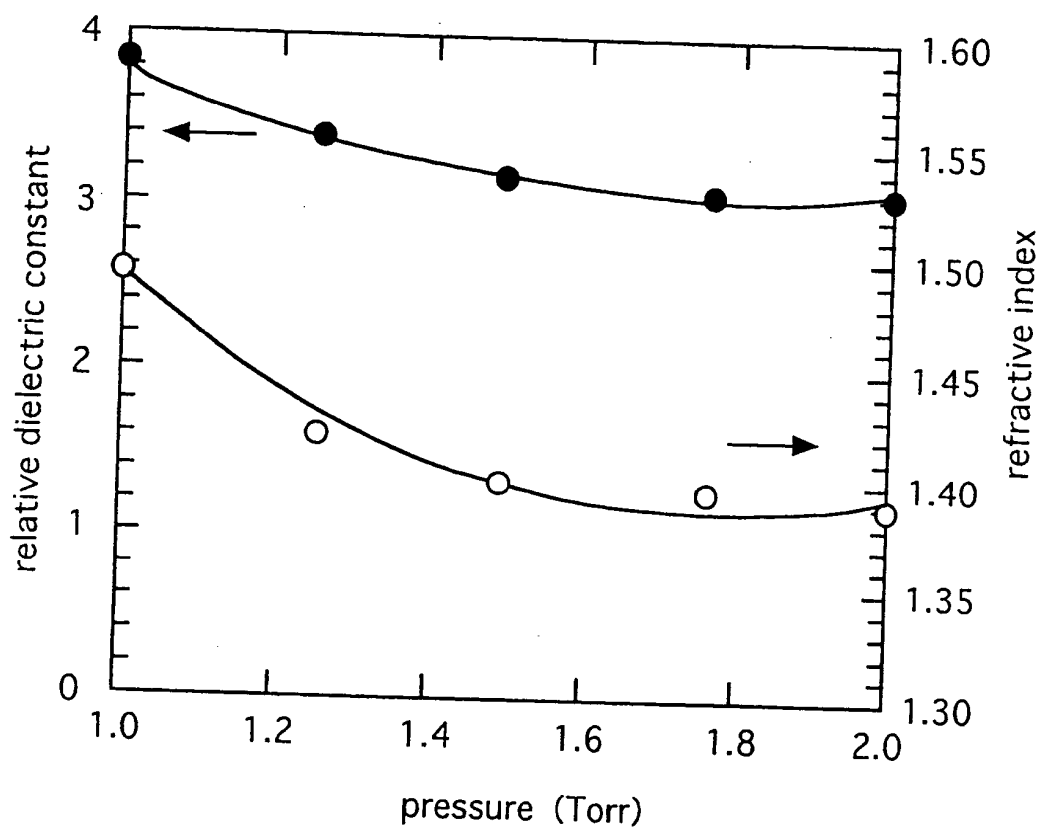
power of the first high-frequency power supply 107 :

300W

power of the second high-frequency power supply 109 :

0W

FIG.6



HMDS flow rate : 50sccm

N₂O flow rate : 200sccm

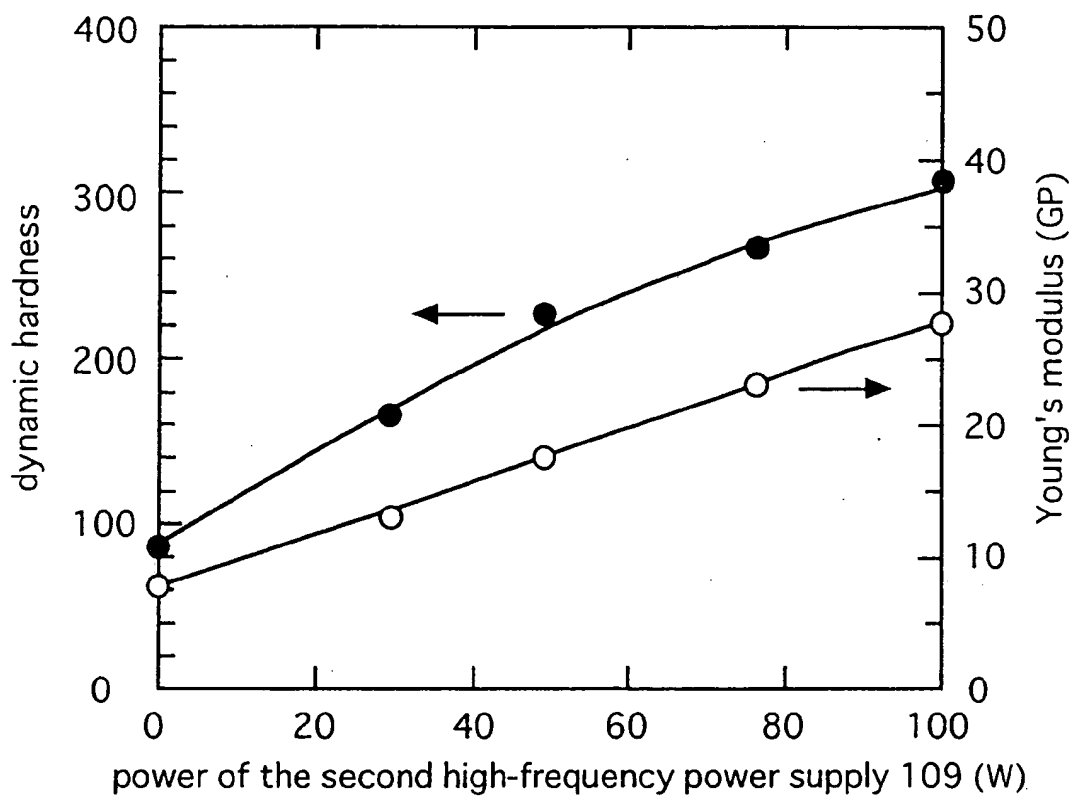
substrate temperature : 350°C

electrode distance : 25mm

power of the first high-frequency power supply 107 :
300W

power of the second high-frequency power supply 109 :
30W

FIG.7



HMDS flow rate : 50sccm

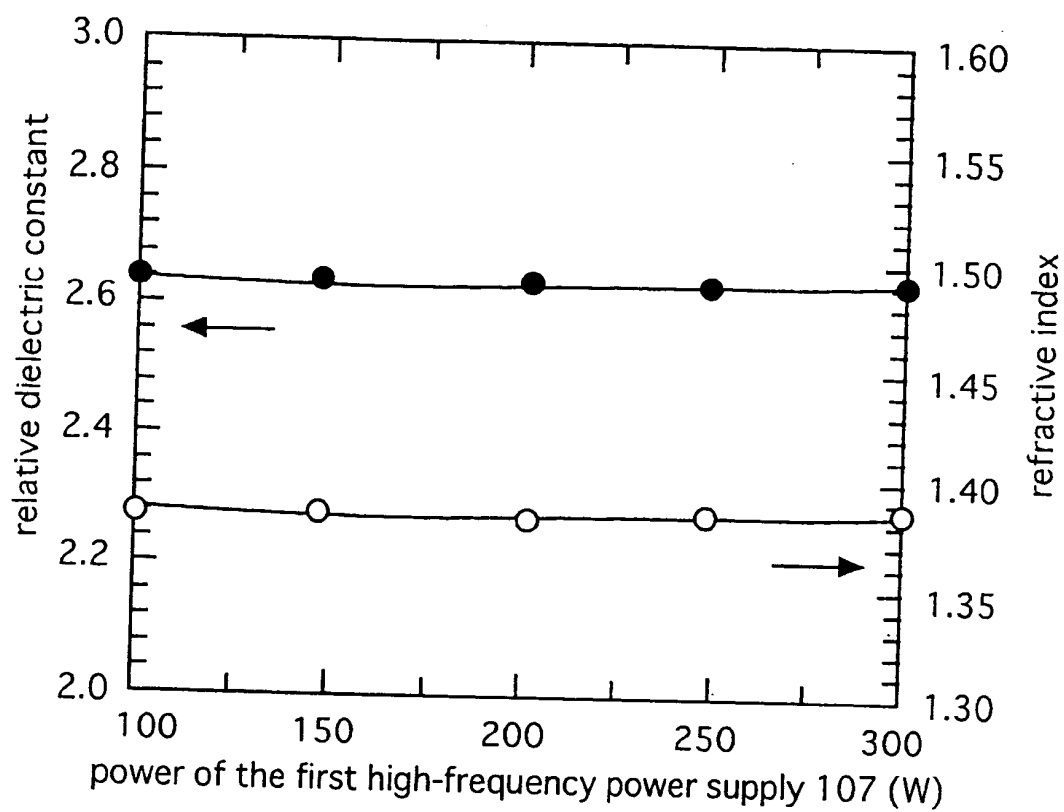
N₂O flow rate : 200sccm

substrate temperature : 350°C

electrode distance : 25mm

power of the first high-frequency power supply 107 :
100W

FIG.8



HMDS flow rate : 50sccm

N₂O flow rate : 200sccm

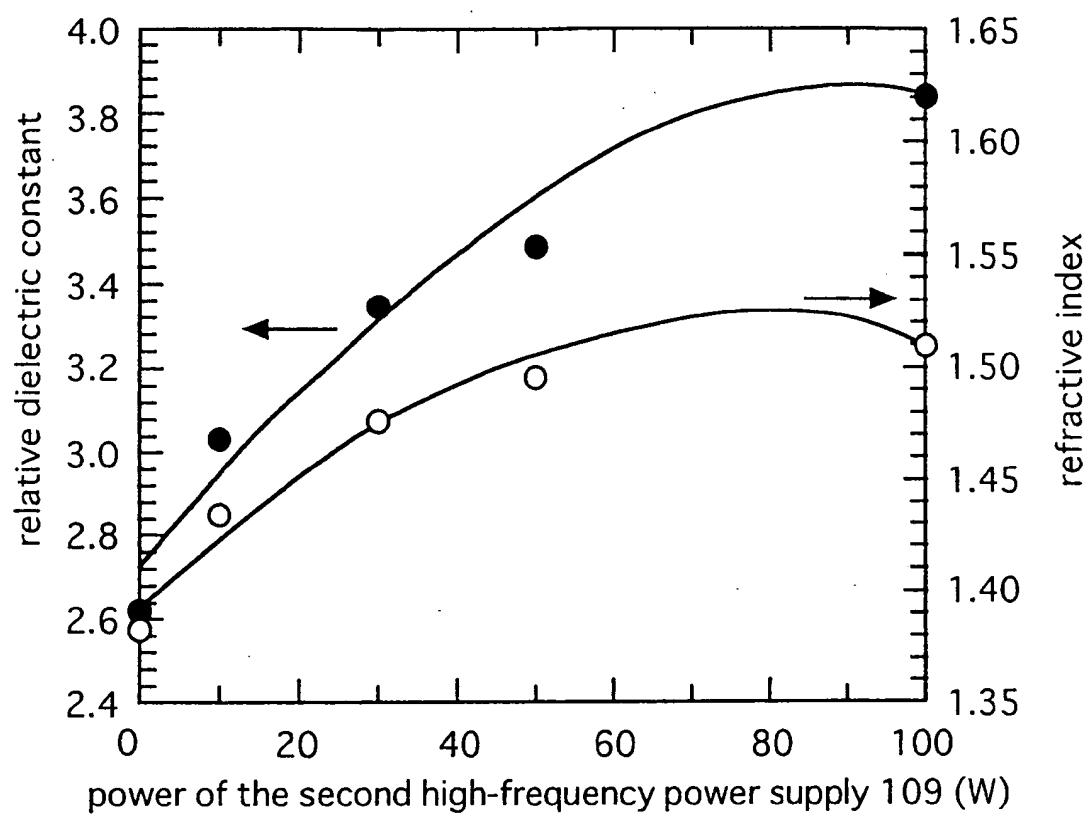
substrate temperature : 350°C

electrode distance : 50mm

pressure : 0.9Torr

power of the second high-frequency power supply 109 :
0W

FIG.9



HMDS flow rate : 50sccm

N₂O flow rate : 200sccm

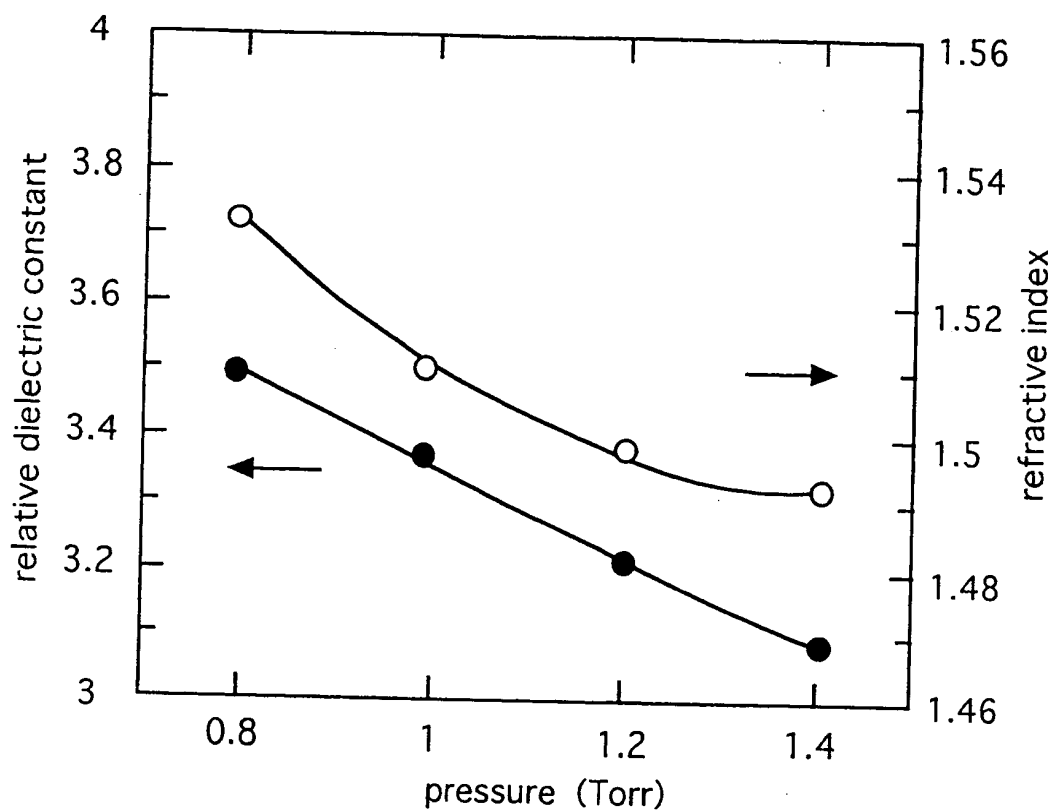
substrate temperature : 350°C

electrode distance : 50mm

pressure : 0.9Torr

power of the first high-frequency power supply 107 :
300W

FIG.10



HMDS flow rate : 50sccm

H₂O flow rate : 100sccm

substrate temperature : 200°C

electrode distance : 25mm

power of the first high-frequency power supply 107 :
0W

power of the second high-frequency power supply 109 :
100W

FIG.11

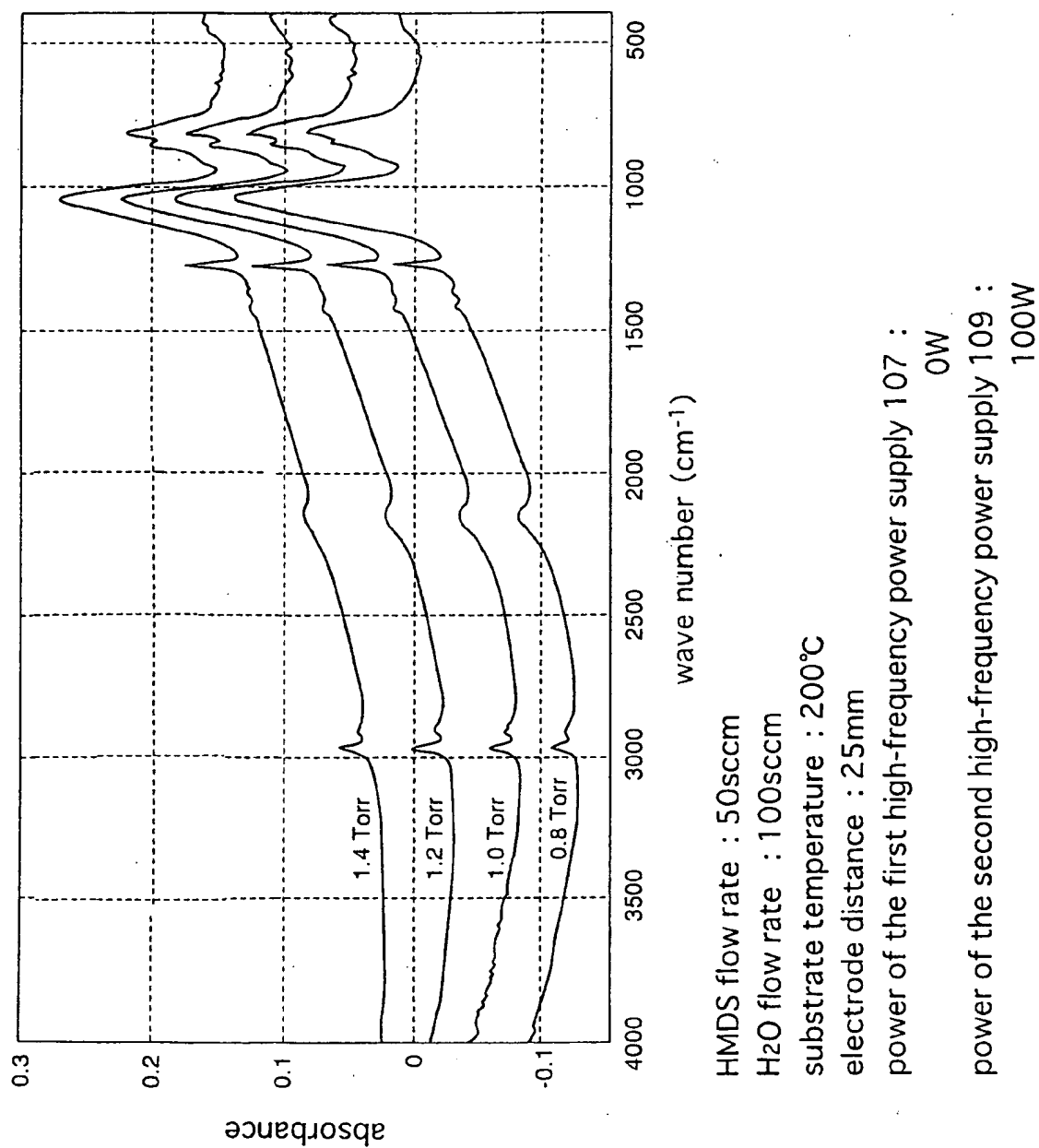
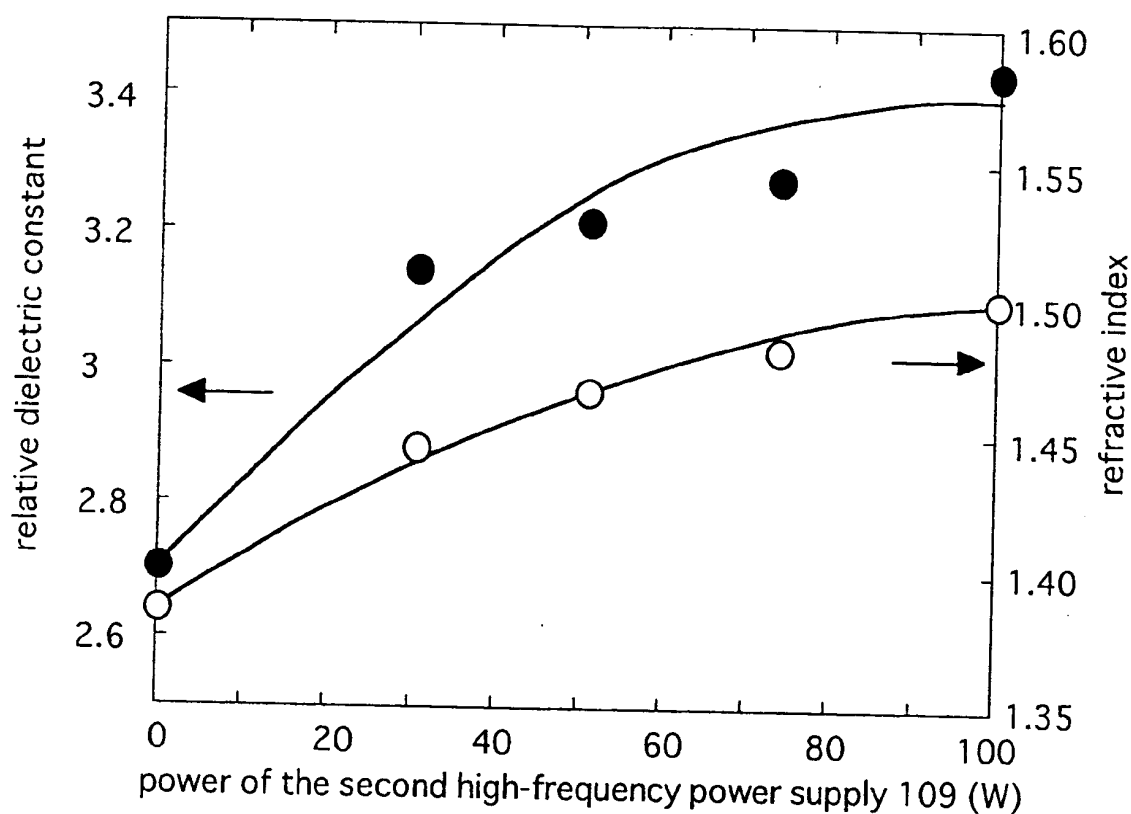


FIG.12



HMDS flow rate : 50sccm

H₂O flow rate : 250sccm

substrate temperature : 375°C

electrode distance : 25mm

pressure : 2.3Torr

power of the first high-frequency power supply 107 :
300W

FIG.13

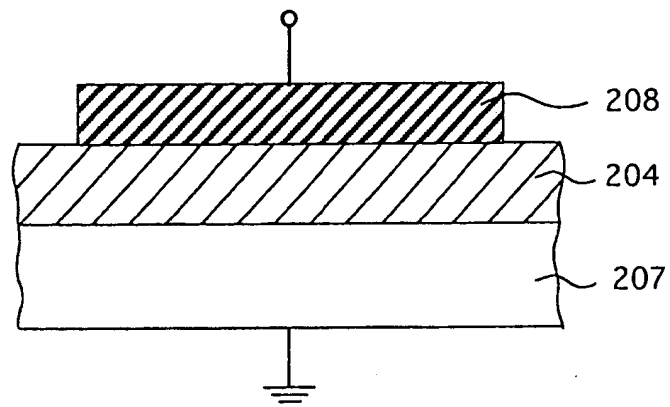
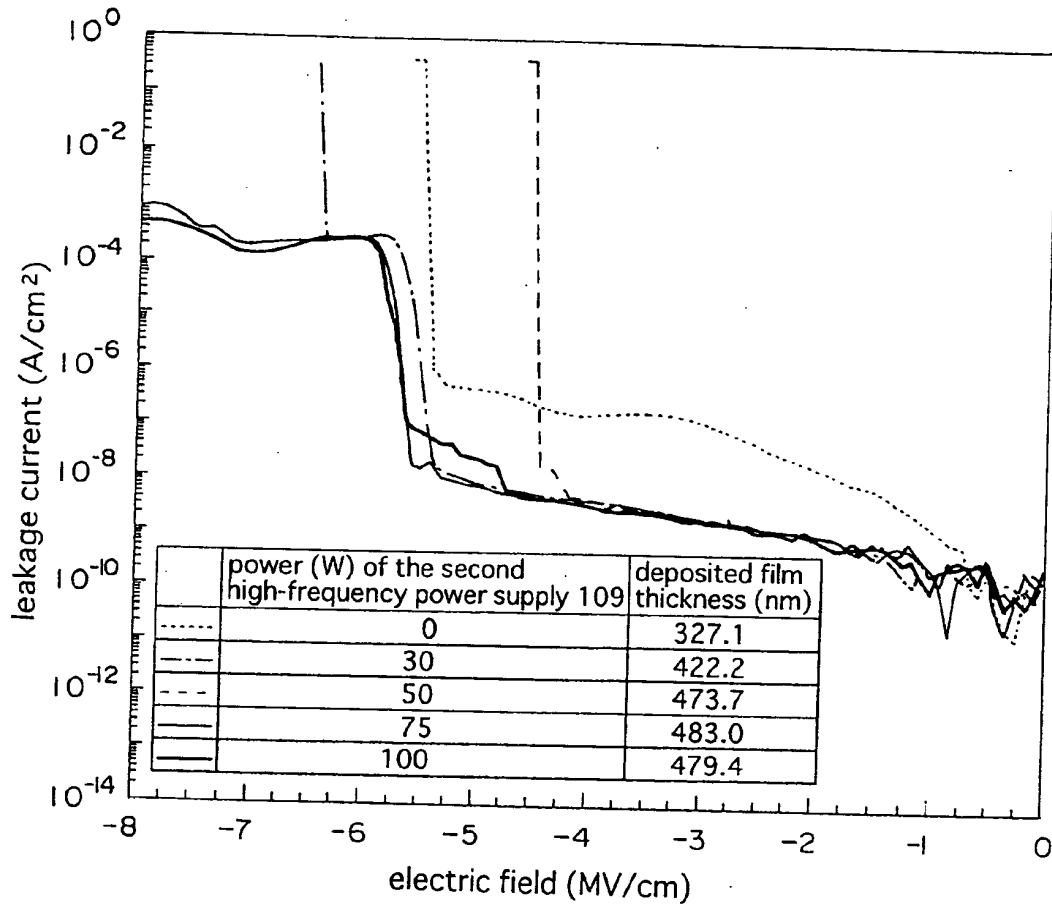


FIG.14



HMDS flow rate : 50sccm

H₂O flow rate : 250sccm

substrate temperature : 375°C

electrode distance : 25mm

pressure : 2.3Torr

power of the first high-frequency power supply 107 :
300W

FIG. 15

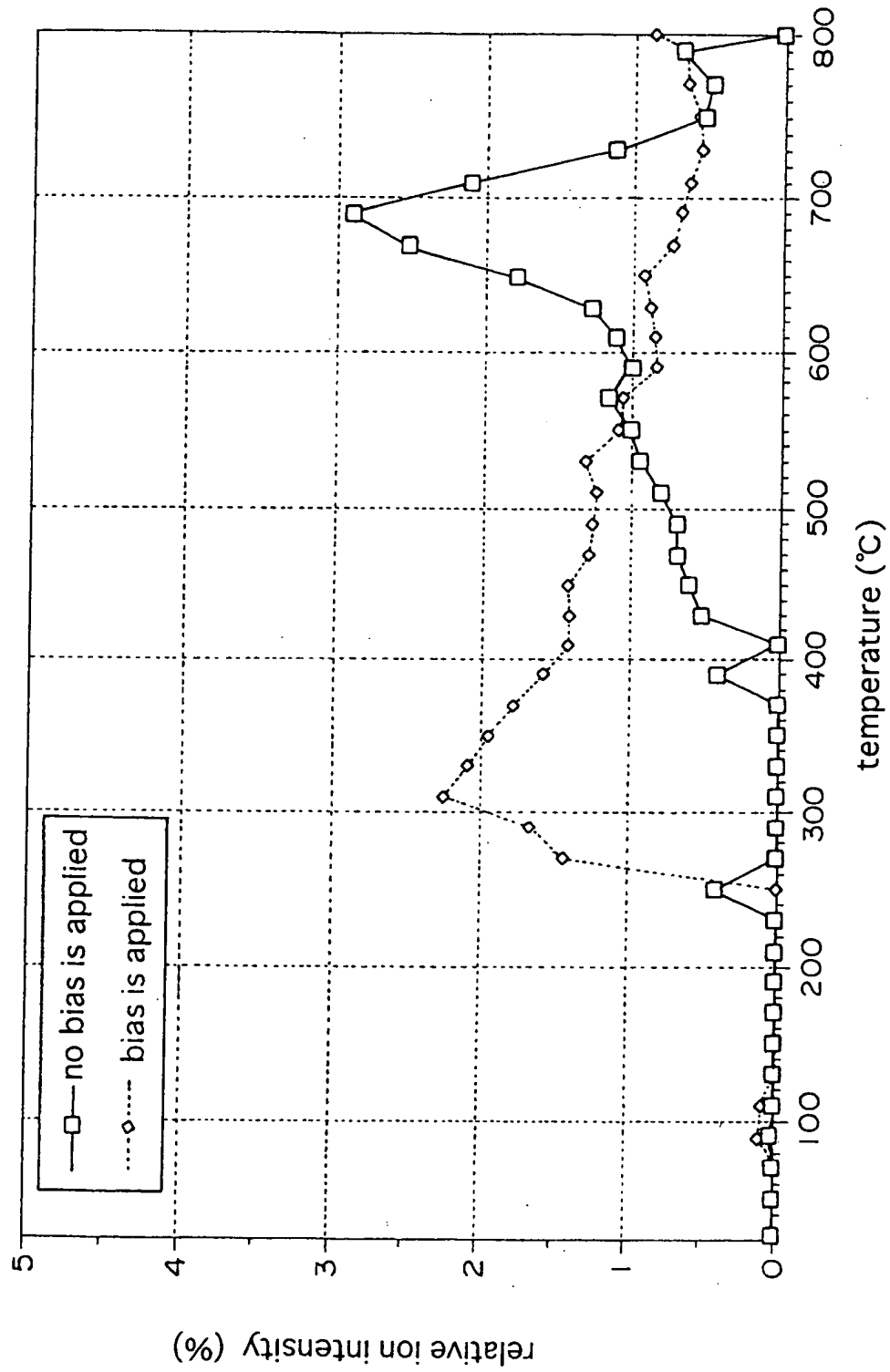


FIG.16

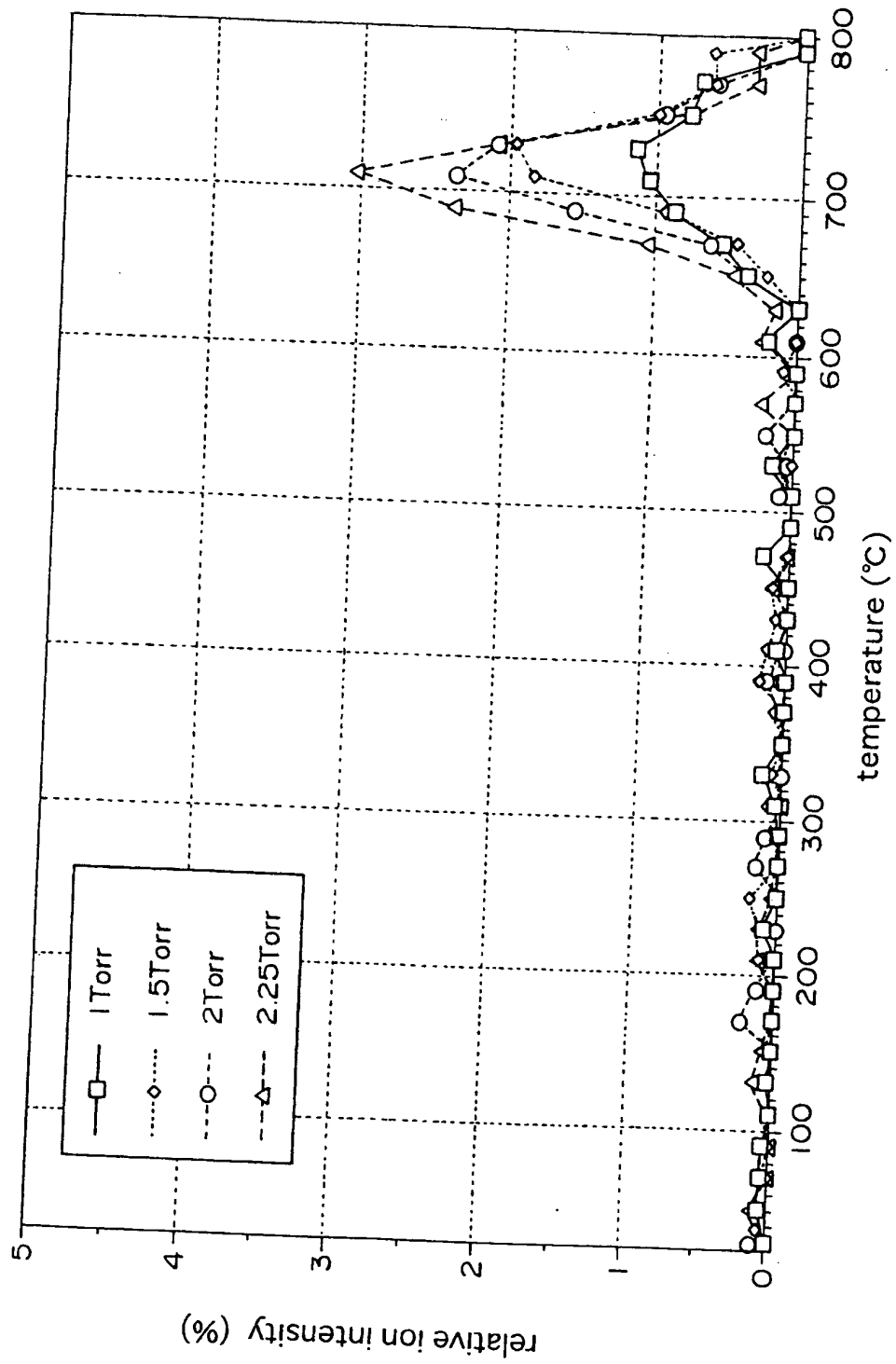
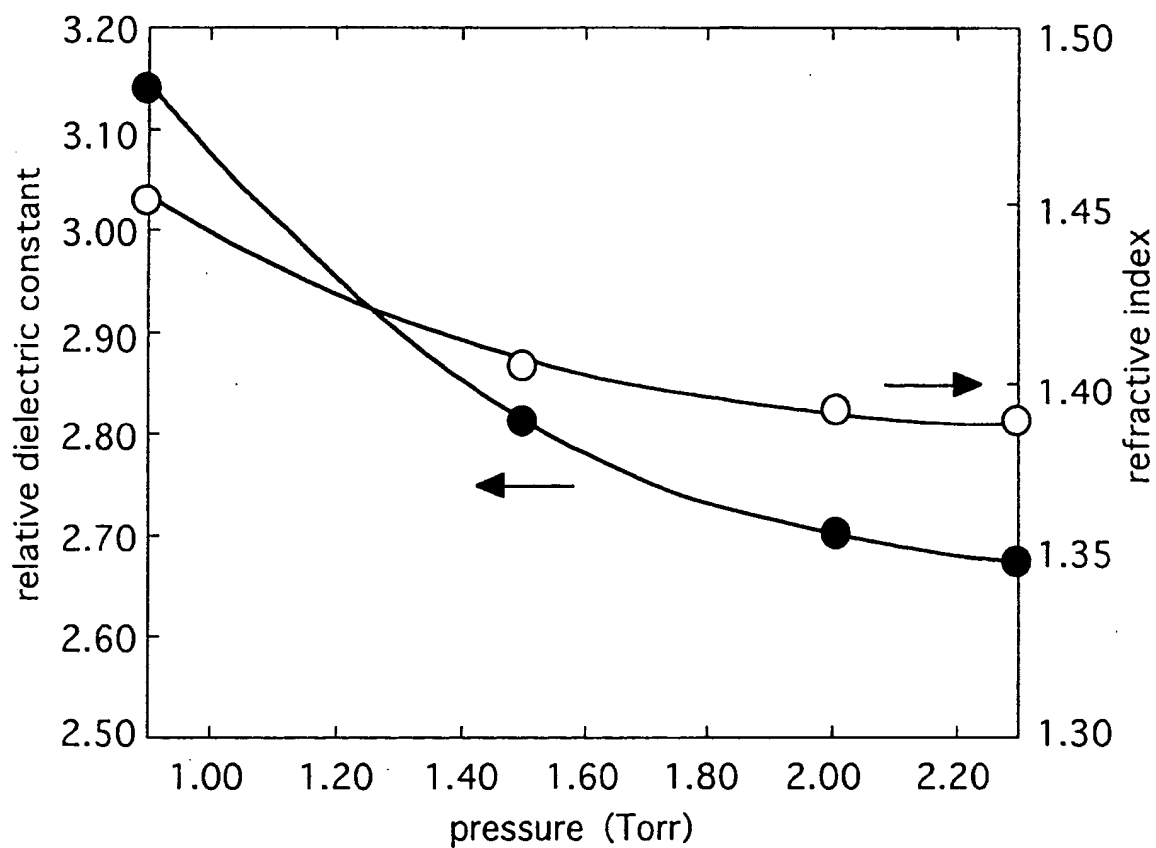


FIG.17



HMDS flow rate : 50sccm

H₂O flow rate : 250sccm

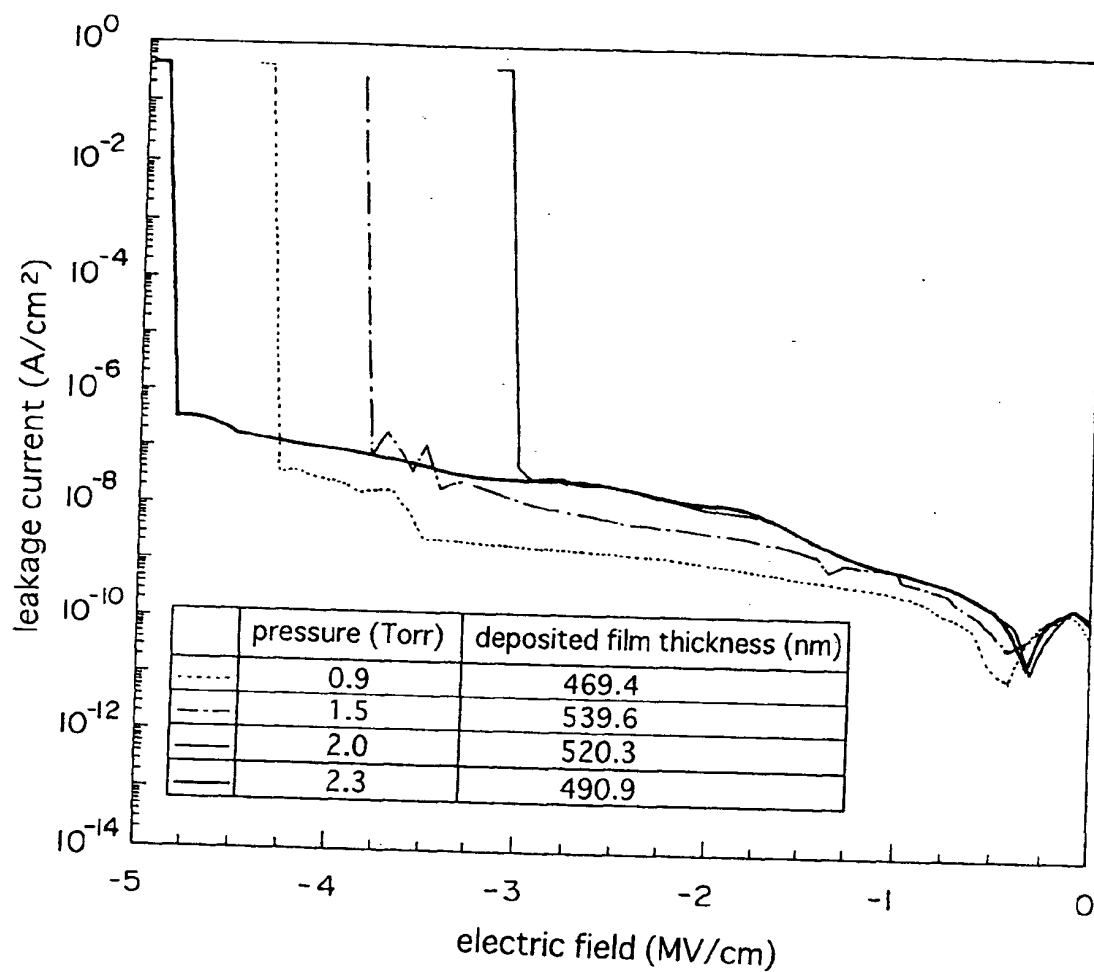
substrate temperature : 375°C

electrode distance : 25mm

power of the first high-frequency power supply 107 :
300W

power of the second high-frequency power supply 109 :
0W

FIG.18



HMDS flow rate : 50sccm

H₂O flow rate : 250sccm

substrate temperature : 375°C

electrode distance : 25mm

power of the first high-frequency power supply 107 :
300W

power of the second high-frequency power supply 109 :
0W

FIG.19A

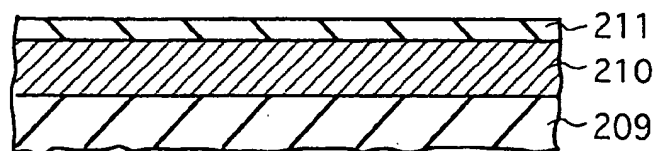


FIG.19B

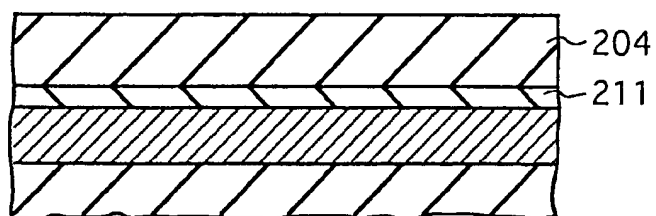


FIG.19C

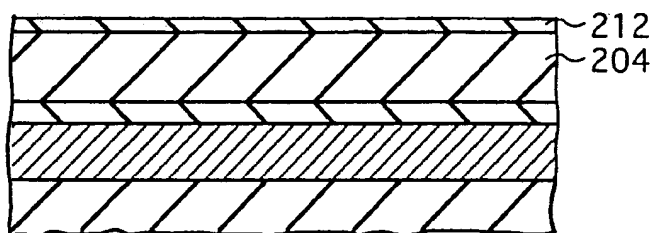


FIG.19D

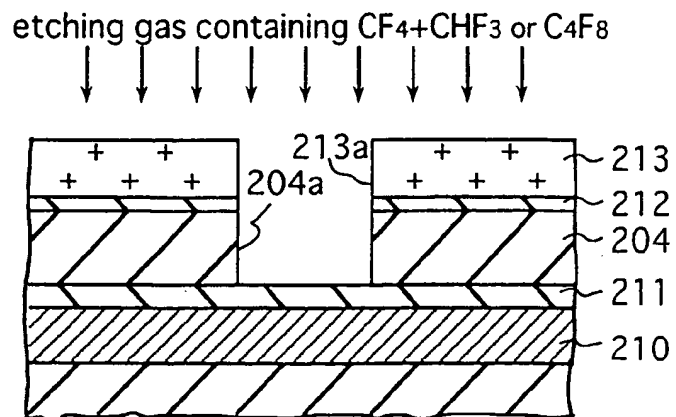


FIG.19E

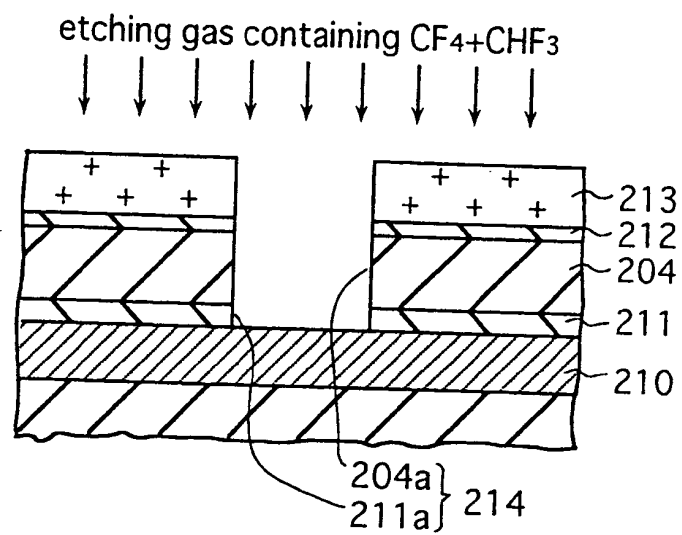


FIG.19F

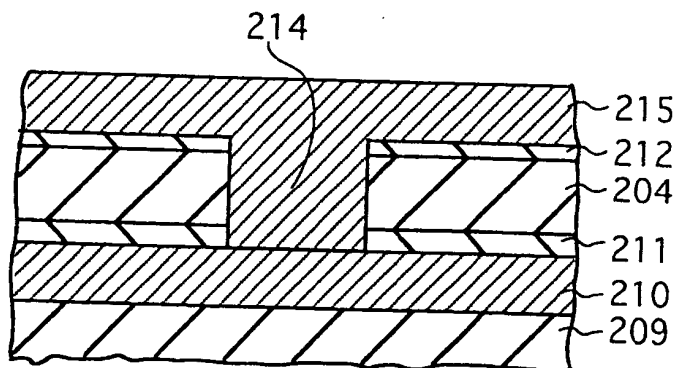


FIG.20A

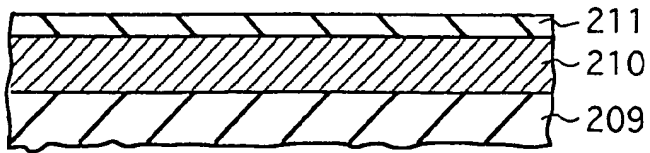


FIG.20B

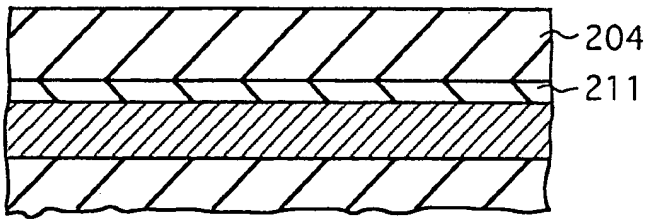


FIG.20C

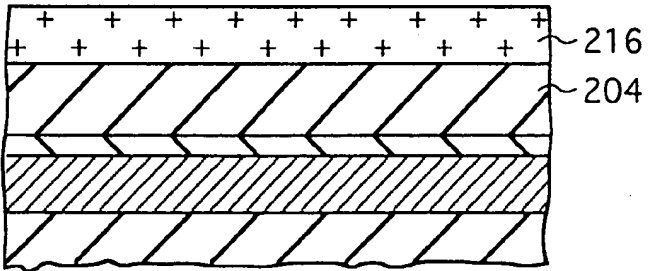


FIG.20D

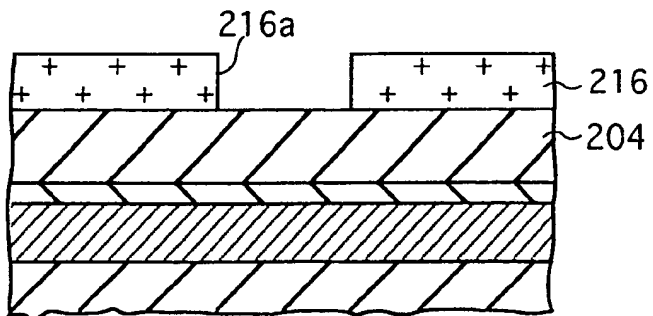


FIG.20E

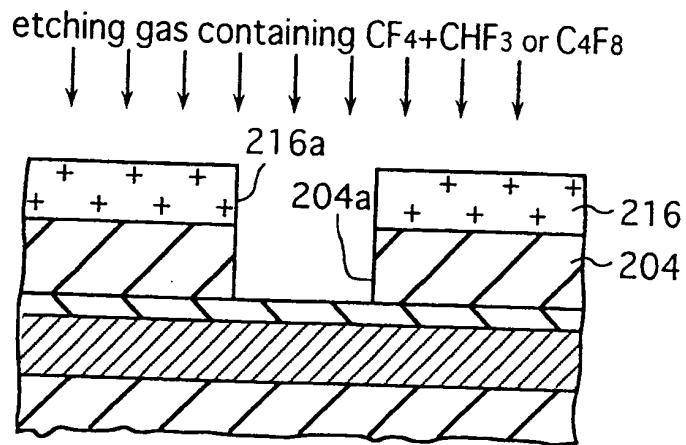


FIG.20F

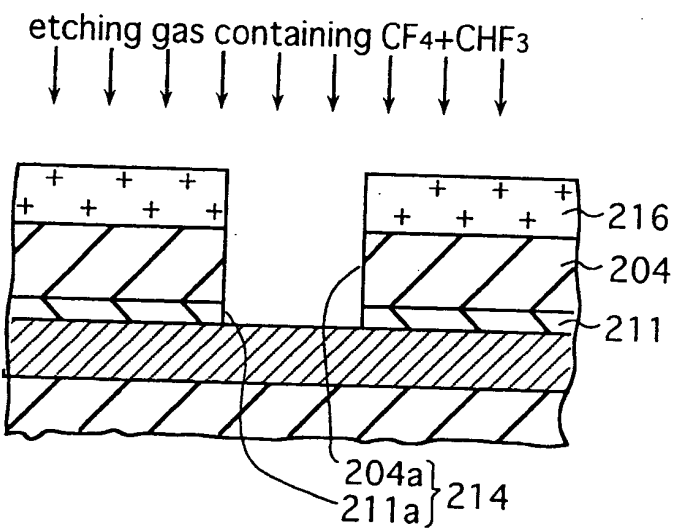
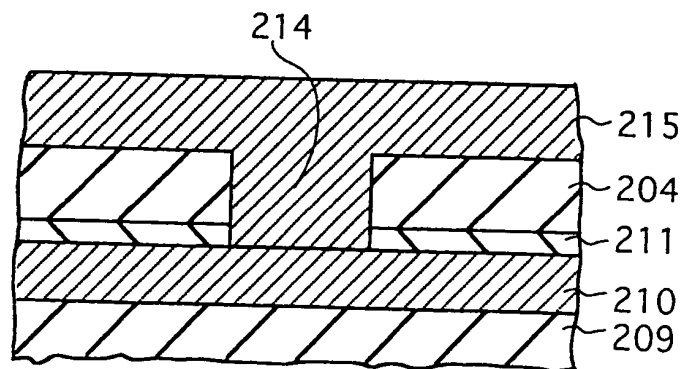


FIG.20G



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